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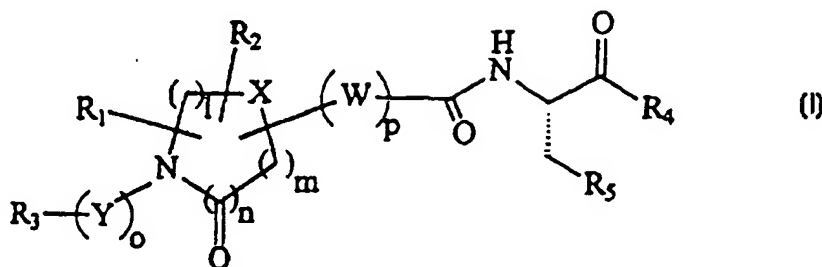
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<p>(21) International Application Number: PCT/US99/14233</p> <p>(22) International Filing Date: 23 June 1999 (23.06.99)</p> <p>(30) Priority Data: 60/090,421 23 June 1998 (23.06.98) US</p> <p>(71) Applicants (for all designated States except US): PHARMACIA & UPJOHN COMPANY [US/US]; 301 Henrietta Street, Kalamazoo, MI 49001 (US). TANABE SEIYAKU CO., LTD. [JP/JP]; 2-10, Dosho-machi 3-chome, Chuo-ku, Osaka 541-8505 (JP).</p> <p>(72) Inventors; and</p> <p>(75) Inventors/Applicants (for US only): BLINN, James, R. [US/US]; 29328 County Road 354, Lawton, MI 49065 (US). CHRUSCIEL, Robert, A. [US/US]; 519 Cherryview Drive, Portage, MI 49024 (US). FISHER, Jed, F. [US/US]; 3517 Old Colony Road, Kalamazoo, MI 49008 (US). TANIS, Steven, P. [US/US]; 7601 Farmington Avenue,</p>		<p>(43) International Publication Date: 29 December 1999 (29.12.99)</p> <p>Kalamazoo, MI 49002 (US). THOMAS, Edward, William [US/US]; 901 Wilshire Boulevard, Kalamazoo, MI 49008 (US). LOBL, Thomas, J. [US/US]; Coulter Pharmaceutical, 600 Gateway Boulevard, South San Francisco, CA 94080 (US). TEEGARDEN, Bradley, R. [US/US]; 11933 Windon Peak Way, San Diego, CA 92131 (US).</p> <p>(74) Agents: MURPHY, Gerald, M., Jr. et al.; Birch, Stewart, Kolasch & Birch, LLP, P.O. Box 747, Falls Church, VA 22042-0747 (US).</p> <p>(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).</p> <p>Published With international search report.</p>

(54) Title: INHIBITORS OF $\alpha_4\beta_1$ MEDIATED CELL ADHESION

(57) Abstract

The present invention relates to compound of formula (I), that are potent inhibitors of $\alpha_4\beta_1$ mediated adhesion to either VCAM or CS-1 and which could be useful for the treatment of inflammatory diseases. Specifically, the molecules of the present invention can be used for treating or preventing $\alpha_4\beta_1$ adhesion mediated conditions in

a mammal such as a human. This method may comprise administering to a mammal or a human patient an effective amount of the compound or composition as explained in the present specification.



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INHIBITORS OF $\alpha_4\beta_1$ MEDIATED CELL ADHESION

Background of the Invention

5

Field of the Invention

The present invention relates to small molecules that are potent inhibitors of $\alpha_4\beta_1$ mediated adhesion to either VCAM or CS-1 and which are useful for the treatment of inflammatory diseases.

10

Description of Related Art

The extracellular matrix (ECM) is the major component of connective tissue which provides structural integrity, and promotes cell migration and differentiation. As part of these functions, extracellular matrix molecules such as fibronectin, collagen, laminin, von
15 Willebrand factor, thrombospondin, fibrinogen, and tenascin have been shown to support adhesion of cells *in vitro*. This adhesive interaction is critical for a number of biological processes including hemostasis, thrombosis, wound healing, tumor metastasis, immunity and inflammation.

Fibronectin (FN) is the prototype ECM molecule. The major cell attachment site in
20 the fibronectin molecule has been reproduced synthetically with the amino acid sequence arginine-glycine-aspartic acid, or RGD using single letter nomenclature. Peptides containing the RGD sequence which either inhibit or promote cell adhesion have been described (US Patent Nos. 4,589,881; 4,661,111; 4,517,686; 4,683,291; 4,578,079; 4,614,517; and 4,792,525). Changes in the peptide as small as the exchange of alanine for glycine or
25 glutamic acid for aspartic acid, which constitute the addition of a single methyl or methylene group to the tripeptide, eliminates these activities (Pierschbacher et al., *Proc. Natl. Acad. Sci. USA* **81**:5985 (1984)). Recently, a second FN cell binding domain has been identified within the alternatively spliced region of the A chain of the molecule, known as the connecting segment 1 (CS-1). The most active cell-binding site within this alternatively spliced region
30 is composed of 25 amino acids where the carboxy terminus contains the sequence

EILDVPST. The amino acid sequence EILDVPST forms a recognition motif on FN for cell surface receptors. (Wayner et al., *J. Cell Biol.* 109:1321 (1989); Guan et al., *Cell* 60:53 (1990)).

5 The receptors which recognize these sites on FN belong to a gene superfamily called integrins which consist of heterodimeric complexes of non-covalently associated alpha and beta subunits. A common β subunit combines with unique I subunits to form an adhesion receptor of defined specificity. To date, 8 β subunits have been identified which can dimerize with 16 distinct I subunits forming 22 distinct integrins. The $\beta 1$ subfamily, also known as the VLA family (Very Late Activation Antigens), binds to ECM molecules such as
10 FN, collagen and laminin. For reviews, see, Hynes, *Cell* 48:549 (1987); Hemler, *Annu. Rev. Immunol.* 8:365 (1990). Leukocyte interaction with FN at the two spatially separate binding domains is mediated by two distinct integrins. The RGD site is recognized by the integrin $\alpha_5\beta_1$, while, EILDV is recognized by $\alpha_4\beta_1$ (Pytela et al., *Cell* 40:191 (1985); Wayner et al., *J. Cell Biol.* 109:1321 (1989); Guan et al, *Cell* 60:53 (1990)).

15 Vascular endothelial cells form the interface between blood and tissues and control the passage of leukocytes as well as plasma fluid into tissues. A variety of signals generated at the site of inflammation can activate both endothelial cells as well as circulating leukocytes so that they become more adhesive to one another. Following this initial adhesion the leukocytes migrate into the tissues to perform host defense functions. Several
20 adhesion molecules have been identified which are involved in leukocyte-endothelial interactions.

In the β_1 subfamily, in addition to binding to fibronectin, $\alpha_4\beta_1$ interacts with a cytokine inducible protein on endothelial cells termed vascular cell adhesion molecule (VCAM). Further involved in the leukocyte-endothelial adhesion process is the β_2 integrin
25 subfamily. β_2 integrins include CD11a/CD18, CD11b/CD18, and CD11c/CD18. In addition, the β_7 subunit associates with α_4 to form a unique $\alpha_4\beta_7$ heterodimer which binds to FN, to VCAM, and to Mucosal Addressin Cell Adhesion Molecule-1 (MAdCAM) (Ruegg et al, *J. Cell.Biol.* 117:179 (1992); Andrew et al., *J. Immunol.* 153:3847 (1994); Briskin et al., *Nature* 363:461 (1993); Shyjan et al, *J. Immunol.* 156:2851 (1996)). α_4 integrins are widely
30 expressed on different cell types including hematopoietic progenitors, lymphocytes, natural

killer cells, monocytes, eosinophils, basophils, and mast cells (Helmer, M. E., *Annu. Rev. Immunol.* 8:365 (1990)). Other molecules on endothelial cells which bind to the leukocytes include ICAM-1, ICAM-2, E-selectin and P-selectin (Carlos and Harlan, *Immunol. Rev.* 114:1 (1990); Osborn, L., *Cell* 62:3 (1990); Springer T., *Nature* 346:425 (1990); Geng et al.,
5 *Nature* 347:757 (1990); Stoolman, *Cell* 56:907 (1989)).

A number of *in vitro* and *in vivo* studies indicate that $\alpha_4\beta_1$ plays a critical role in the pathogenesis of a variety of diseases. Monoclonal antibodies directed against α_4 have been tested in a variety of disease models. Anti- α_4 antibodies block adhesion of lymphocytes to synovial endothelial cells; this adhesion plays a potential role in rheumatoid arthritis (van
10 Dinther-Janssen et al, *J. Immunol.* 147:4207 (1991)). α_4 has also been implicated with respect to rheumatoid arthritis in separate studies (Laffon et al, *J. Clin. Invest.* 88:546 (1991); Morales-Ducet et al, *J. Immunol.* 149:1424 (1992)). A significant number of studies have evaluated the role of α_4 in allergy and asthma. For example, monoclonal antibodies to α_4 block adhesion of basophils and eosinophils to cytokine activated endothelial cells (Walsh et
15 al, *J. Immunol.* 146:3419 (1991); Bochner et al, *J. Exp. Med.* 173:1553 (1991)). Monoclonal antibodies to α_4 were also effective in several lung antigen challenge models (Abraham et al, *J. Clin. Invest.* 93:776 (1994); Weg et al, *J. Exp. Med.* 177:561 (1993)). The cotton-top tamarin, which experiences spontaneous chronic colitis, showed a significant attenuation of their colitis when anti- α_4 antibody was administered (Podolsky et al, *J. Clin.*
20 *Invest.* 92:372 (1993); Bell et al, *J. Immunol.* 151:4790 (1993)). In a rat and mouse model, autoimmune encephalomyelitis was blocked by anti- α_4 antibody (Yednock et al, *Nature* 356:63 (1992); Baron et al, *J. Exp. Med.* 177:57 (1993)). Anti- α_4 monoclonal antibodies also inhibit insulinitis and delay the onset of diabetes in the non-obese diabetic mouse (Baron et al, *J. Clin. Invest.* 93:1700 (1994); Yang et al, *Proc. Natl. Acad. Sci. USA* 90:10494
25 (1993); Burkly et al, *Diabetes* 43:529 (1994)). α_4 is also implicated in atherosclerosis due to its endothelial expression during atherogenesis (Cybulsky et al, *Science* 251:788 (1991)). The migration of leukocytes to an inflammatory site can also be blocked by anti- α_4 antibodies. In addition to the blocking of migration, inhibitors of leukocyte endothelial adhesion may block the costimulatory signals mediated by integrins and thus inhibit
30 overproduction of inflammatory cytokines. In a separate set of experiments not using anti- α_4

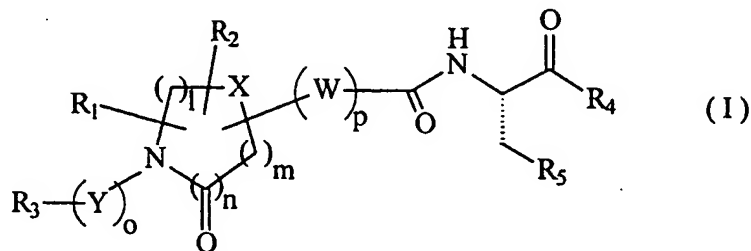
antibodies, the peptides GRDGSP or EILDV were tested against contact hypersensitivity response. The contact hypersensitivity response was found to be blocked by GRDGSP or EILDV suggesting that both $\alpha_4\beta_1$ and $\alpha_5\beta_1$ are involved in this inflammatory response.

Other ailments which may involve $\alpha_4\beta_1$ -mediated conditions include the inflammatory disorders rheumatoid arthritis, allergic disorders, asthma, spontaneous chronic colitis, insulinitis, contact hypersensitivity response, atherosclerosis and autoimmune encephalomyelitis. These studies illustrate that small molecules that are potent inhibitors of $\alpha_4\beta_1$ mediated adhesion to either VCAM-1 or CS-1 may be used as a form of treatment in numerous inflammatory diseases. However, these inflammatory conditions could be expanded to include adult respiratory distress syndrome, AIDS, cardiovascular diseases, thrombosis or harmful platelet aggregation, reocclusion following thrombolysis, allograft rejection, reperfusion injury, psoriasis, eczema, contact dermatitis and other skin inflammatory diseases, osteoporosis, osteoarthritis, atherosclerosis, neoplastic diseases including metastasis of neoplastic or cancerous growth, wound healing enhancement, treatment of certain eye diseases such as detaching retina, Type I diabetes, multiple sclerosis, systemic lupus erythematosus (SLE), inflammatory and immunoinflammatory conditions including ophthalmic inflammatory conditions and inflammatory bowel diseases, ulcerative colitis, regional enteritis and other autoimmune diseases. Accordingly, a compound which could inhibit these conditions is desirable.

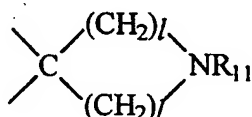
Summary of the Invention

The present invention particularly provides:

A compound of the formula:

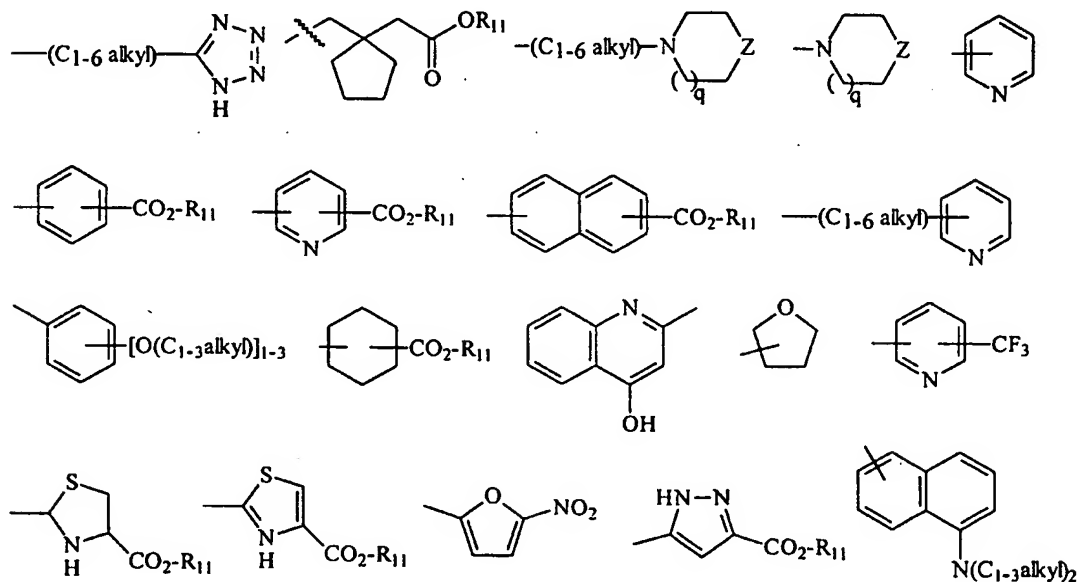


In the above formula (I), R_1 may occur one to four times and each occurrence is independently hydrogen or C_{1-6} alkyl. Also in the above formula (I), R_2 is hydrogen, pyridyl, C_{1-6} alkyl, $(C_{1-6} \text{ alkyl})\text{-CO}_2\text{-R}_{11}$, or $\text{-CO}_2\text{-R}_{11}$. In addition, R_1 and R_2 may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



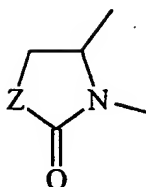
together with the carbon atom to which they are attached.

In the above formula (I), R_3 is hydrogen, phenyl, C_{1-6} alkyl, C_{3-6} alkenyl, C_{7-17} arylalkyl, $(C_{1-6} \text{ alkyl})\text{-CO}_2\text{-R}_{11}$, $(C_{2-6} \text{ alkenyl})\text{-CO}_2\text{-R}_{11}$, $(C_{1-6} \text{ alkyl})\text{-CO-C}_{1-6} \text{ alkyl}$, $(C_{1-6} \text{ alkyl})\text{-O-C}_{1-6} \text{ alkyl}$, $(C_{1-6} \text{ alkyl})\text{-OH}$, $(C_{1-6} \text{ alkyl})\text{-CN}$, adamantyl or one of the following:

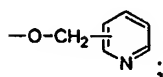


In addition, R_2 and $R_3\text{-(Y)}_0\text{-}$ may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:

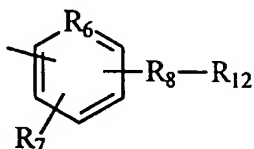
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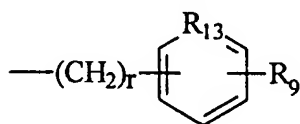
In the above formula (I), R_4 is $-O-R_{11}$, NH_2 , $NHOH$, $-O-(C_{7-10} \text{ arylalkyl})$, or is of the formula:



In the above formula (I), R_5 is a formula of the following:



- 5 In the above, R_6 is N or CH, R_7 is hydrogen or halogen, R_8 is $-NH-Y_1-$, $-OCH_2-$, or $-CONH-$, R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or trifluoromethyl, R_{10} is C_{1-6} alkyl, or $(C_{1-6} \text{ alkyl})-OH$, or hydrogen, R_{11} is hydrogen or C_{1-6} alkyl, R_{12} is C_{1-6} alkyl or the following formula:



, and R_{13} is N or CH.

- 10 In the above formula (I), W is $(C_{1-6} \text{ alkyl})$, X is S, O, or CH_2 , Y and Y_1 are independently $-CO-$, $-C(=O)O-$, $-SO_2-$, or $-C(=O)N(R_{10})-$ and Z is O, CH_2 , or $N-R_{11}$.

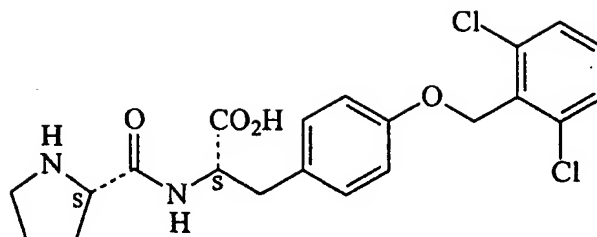
In the above, l is 1, 2, or 3, m is 1 or 2, n is 0 or 1, o is 0 or 1, p is 0 or 1, q is 0 or 1, and r is 0, 1, 2 or 3.

The above formula (I) has the provisos that:

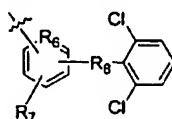
- 15 (1) when Y is $-C(=O)O-$, R_3 cannot be hydrogen;
 (2) when R_4 is equal to $O-(C_4 \text{ alkyl})$, C_4 alkyl is not equal to *tert*-butyl;
 (3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is CH_2), W is equal to CH_2 ; and

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(4) the compound is not



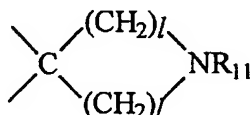
In another embodiment of the present invention, R_5 is a formula of the following:



wherein R_7 is hydrogen or Cl.

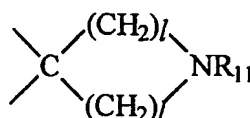
In another embodiment of the present invention, R_2 is hydrogen or C_{1-6} alkyl.

Additionally, R_1 and R_2 may be attached to the same carbon atom and may form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached or
 5 be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



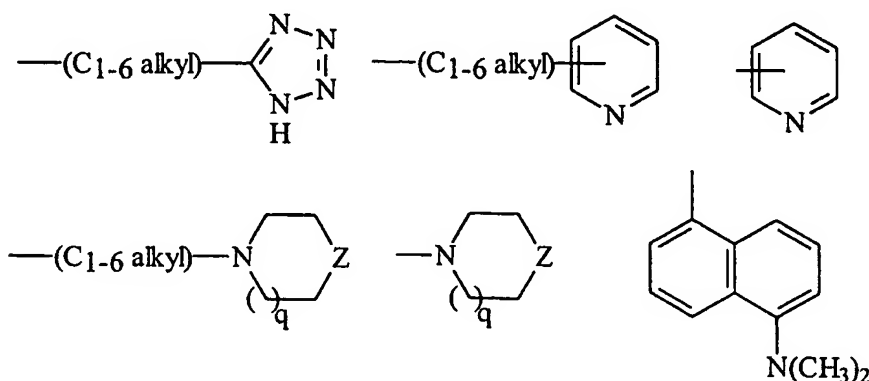
together with the carbon atom to which they are attached. Also, in this embodiment, n is 0, m is 2 and p is 0.

10 In yet another embodiment of the present invention, R_1 is hydrogen or C_{1-3} alkyl, and R_2 is hydrogen or C_{1-4} alkyl. Additionally, R_1 and R_2 may be attached to the same carbon atom and may form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached or be attached to the same atom and form a ring of 5-8 atoms of the formula:

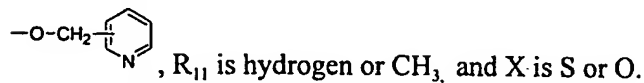


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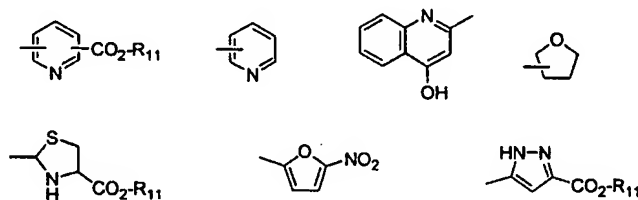
together with the carbon atom to which they are attached. Also in this embodiment, R_3 is hydrogen, C_{1-6} alkyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)-OH, $(C_{1-6}$ alkyl)-CO₂-R₁₁, $(C_{1-6}$ alkyl)-CN, adamantyl, phenyl, or one of the following:



Additionally, in this embodiment, R_4 is -O-R₁₁, NH₂, NHOH, or R_4 is of the formula:



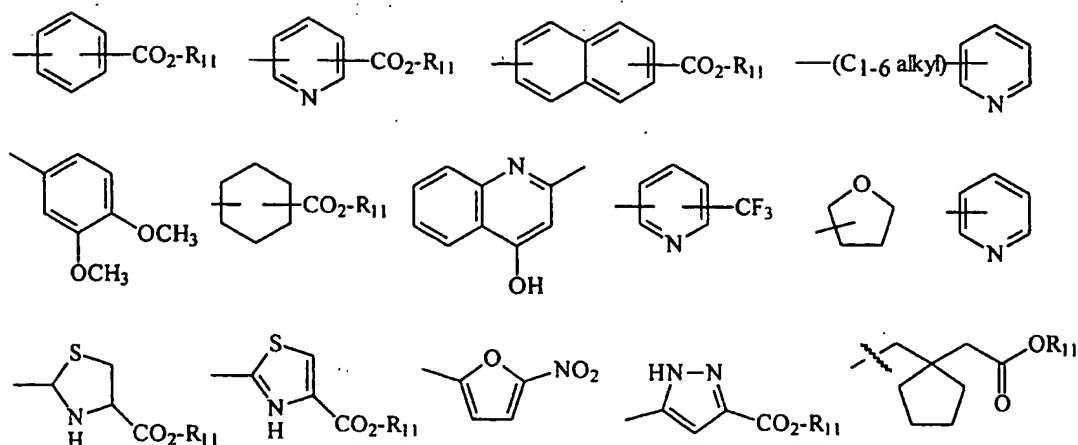
- 5 In another embodiment, R_1 and R_2 are hydrogen, and R_3 is $(C_{2-6}$ alkenyl)-CO₂-R₁₁, $(C_{1-6}$ alkyl)-O-C₁₋₃ alkyl, $(C_{1-6}$ alkyl)-CO₂-R₁₁, or one of the following:



Also, in this embodiment, R_4 is O-R₁₁, R_6 is CH, R_{11} is hydrogen, and R_7 is hydrogen.

Additionally, X is S, Y is -CO-, and l is 1.

- 10 In another embodiment of the present invention, R_1 and R_2 are hydrogen, and R_3 is C_{1-6} alkyl, $(C_{1-6}$ alkyl)-CO₂-R₁₁, $(C_{2-6}$ alkenyl)-CO₂-R₁₁, $(C_{1-6}$ alkyl)-CO-C₁₋₆ alkyl, $(C_{1-6}$ alkyl)-O-C₁₋₃ alkyl, $(C_{1-6}$ alkyl)-CN, or one of the following:

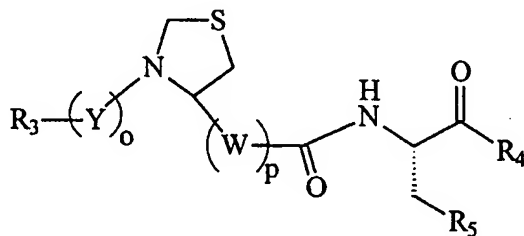


Additionally, R_4 is OH, R_6 is CH, R_{11} is hydrogen, R_7 is hydrogen, X is CH_2 , and Y is CO- or -C(=O)NH- .

In another embodiment of the present invention, W is $(\text{C}_{1-3} \text{ alkyl})$, X is CH_2 , Y is -C(=O)O- , R_1 is hydrogen, R_2 is hydrogen, $(\text{C}_{1-3} \text{ alkyl})\text{-CO}_2\text{-R}_{11}$, or $\text{-CO}_2\text{-R}_{11}$, R_3 is hydrogen, C_{7-10} arylalkyl, C_{1-6} alkyl, or $(\text{C}_{1-6} \text{ alkyl})\text{-CO}_2\text{-R}_{11}$, R_4 is OH, R_6 is CH, R_{11} is hydrogen, R_7 is hydrogen, l is 1 or 3, and n is 0.

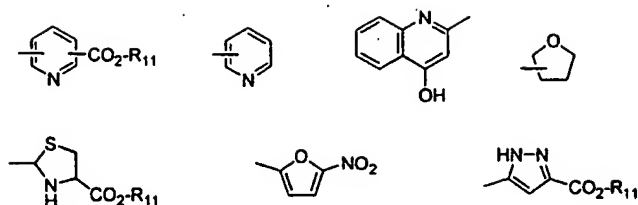
In another embodiment of the present invention, W is C_{1-3} alkyl, X is CH_2 , R_1 is hydrogen, R_2 is $(\text{C}_{1-4} \text{ alkyl})\text{-CO}_2\text{-R}_{11}$, or $\text{CO}_2\text{-R}_{11}$, R_3 is hydrogen, C_{1-3} alkyl, or C_{2-7} alkenyl, R_4 is OH, R_6 is CH, R_{11} is hydrogen, R_7 is hydrogen, l is 1, m is 1, n is 1, o is 0, and p is 1.

In another embodiment, the compound of the present invention is represented by the following formula



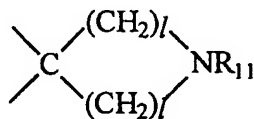
In another embodiment of the present invention, R_1 and R_2 are hydrogen, and R_3 is C_{1-6} alkyl, $(\text{C}_{1-6} \text{ alkyl})\text{-CO}_2\text{R}_{11}$, $(\text{C}_{2-6} \text{ alkenyl})\text{-CO}_2\text{R}_{11}$, $(\text{C}_{1-6} \text{ alkyl})\text{-O-C}_{1-3} \text{ alkyl}$, or one of the following:

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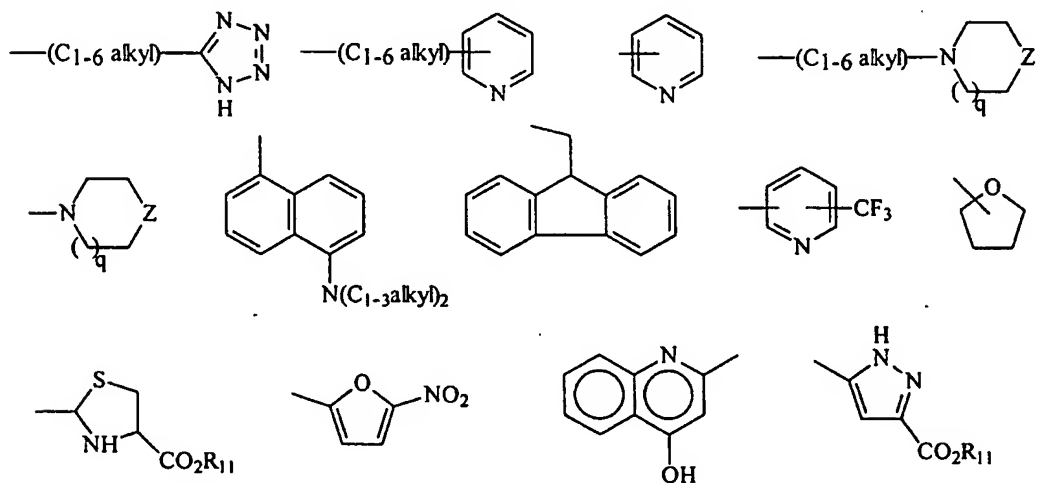


Additionally, in this embodiment, R_4 is $O-R_{11}$, R_6 is CH , R_{11} is hydrogen or C_{1-6} alkyl, R_7 is hydrogen, X is S , Y is $-C(=O)O-$, and l is 1.

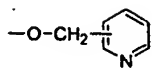
In another embodiment of the present invention, R_1 is hydrogen or C_{1-3} alkyl and R_2 is hydrogen or C_{1-4} alkyl. Additionally, R_1 and R_2 may be attached to the same carbon atom and may form a carbocyclic ring of 5-8 atoms, or be attached to the same atom and form a ring of 5-8 atoms of the formula:



together with the carbon atom to which they are attached. In this embodiment, R_3 is hydrogen, C_{1-6} alkyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)-OH, $(C_{1-6}$ alkyl)- CO_2-R_{11} , $(C_{1-6}$ alkyl)-CN, adamantyl, phenyl, or one of the following:

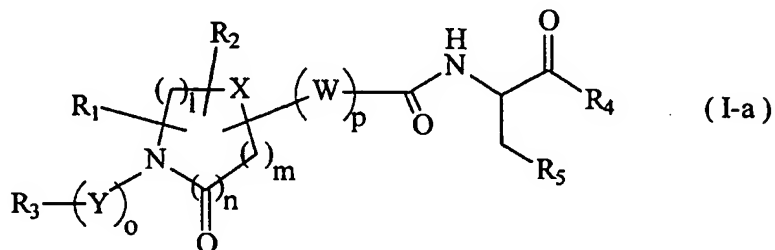


Additionally, in this embodiment, R_4 is $-O-R_{11}$, NH_2 , $NHOH$, or R_4 is of the formula

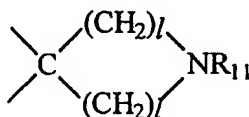


. Also, R₁₁ is hydrogen or C₁₋₆ alkyl, and X is S or O.

In yet another embodiment, the compound of the present invention is represented by the following formula (I-a):



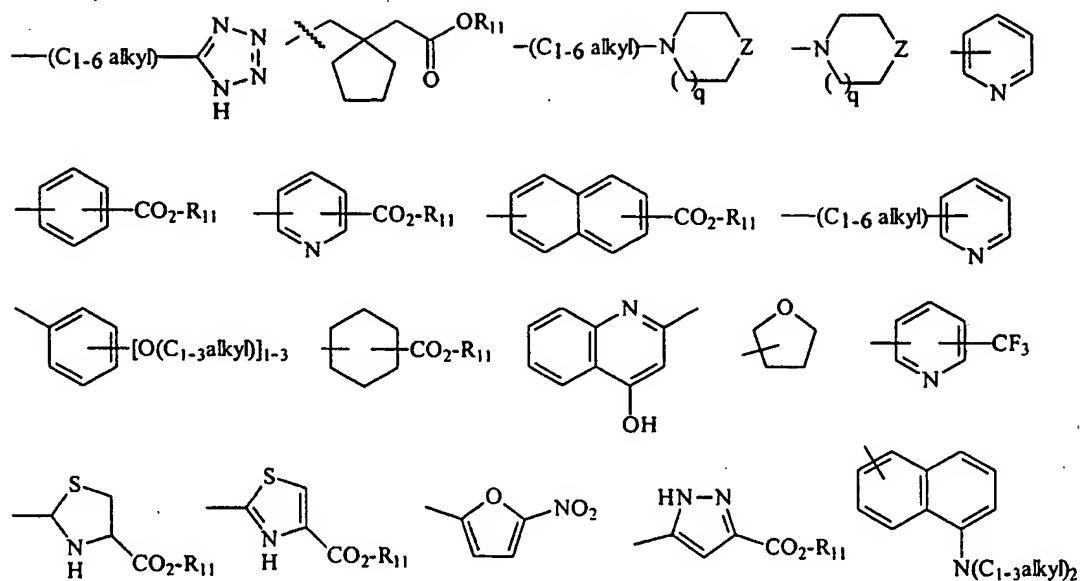
In the above formula (I-a), R₁ may occur one to four times and each occurrence is independently hydrogen or C₁₋₆ alkyl, and R₂ is hydrogen, pyridyl, C₁₋₆ alkyl, (C₁₋₆ alkyl)-CO₂-R₁₁, or -CO₂-R₁₁. Additionally, R₁ and R₂ may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



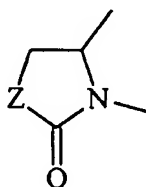
10

together with the carbon atom to which they are attached. Additionally, in this embodiment, R₃ is hydrogen, phenyl, C₁₋₆ alkyl, C₃₋₆ alkenyl, C₇₋₁₇ arylalkyl, (C₁₋₆ alkyl)-CO₂-R₁₁, (C₂₋₆ alkenyl)-CO₂-R₁₁, (C₁₋₆ alkyl)-CO-C₁₋₆ alkyl, (C₁₋₆ alkyl)-O-C₁₋₆ alkyl, (C₁₋₆ alkyl)-OH, (C₁₋₆ alkyl)-CN, adamantyl or one of the following:

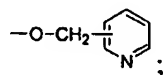
-12-



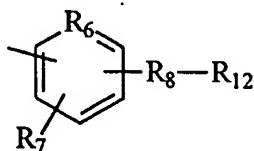
In addition, R_2 and $R_3-(Y)_o$ may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:



- 5 Additionally, R_4 is $-O-R_{11}$, NH_2 , $NHOH$, $-O-(C_{7-10} \text{ arylalkyl})$, or is of the formula



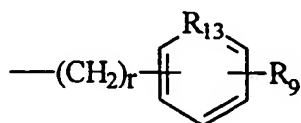
Also, in this embodiment, R_5 is a formula of the following:



Additionally, in this embodiment, R_6 is N or CH, R_7 is hydrogen or halogen, R_8 is

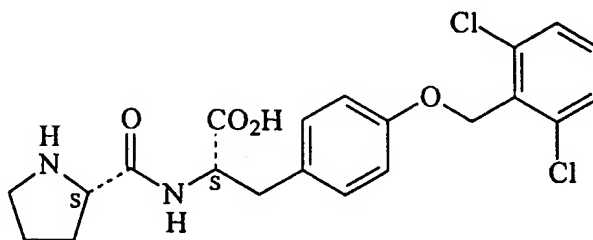
-13-

-NH-Y₁-, -OCH₂-, or -CONH-, R₉ may occur one to three times and is a halogen, C₁₋₆ alkoxy, C₁₋₆ alkyl or trifluoromethyl, R₁₀ is C₁₋₆ alkyl, or (C₁₋₆ alkyl)-OH, or hydrogen, R₁₁ is hydrogen or C₁₋₆ alkyl, R₁₂ is C₁₋₆ alkyl or the following formula:



, R₁₃ is N or CH, W is (C₁₋₆ alkyl), X is S, O, or CH₂, Y and Y₁ are independently -CO-, -C(=O)O-, -SO₂-, or -C(=O)N(R₁₀)-, Z is O, CH₂, or N-R₁₁, *l* is 1, 2, or 3, *m* is 1 or 2, *n* is 0 or 1, *o* is 0 or 1, *p* is 0 or 1, *q* is 0 or 1 and *r* is 0, 1, 2 or 3. This particular embodiment of the present invention has the provisos that

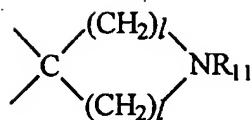
- (1) when Y is -C(=O)O-, R₃ cannot be hydrogen;
- (2) when R₄ is equal to O-(C₄ alkyl), C₄ alkyl is not equal to *tert*-butyl;
- (3) in those pyrrolidine structures (*l* is 1; *m* is 2; *n* is 0; *o* is 0; *p* is 1; X is CH₂), W is equal to CH₂;
- (4) the compound has an IC₅₀ value of less than 5μM in a Jurkat CS-1 assay and/or an IC₅₀ value of less than 50μM in a Jurkat EC assay; and
- (5) the compound is not



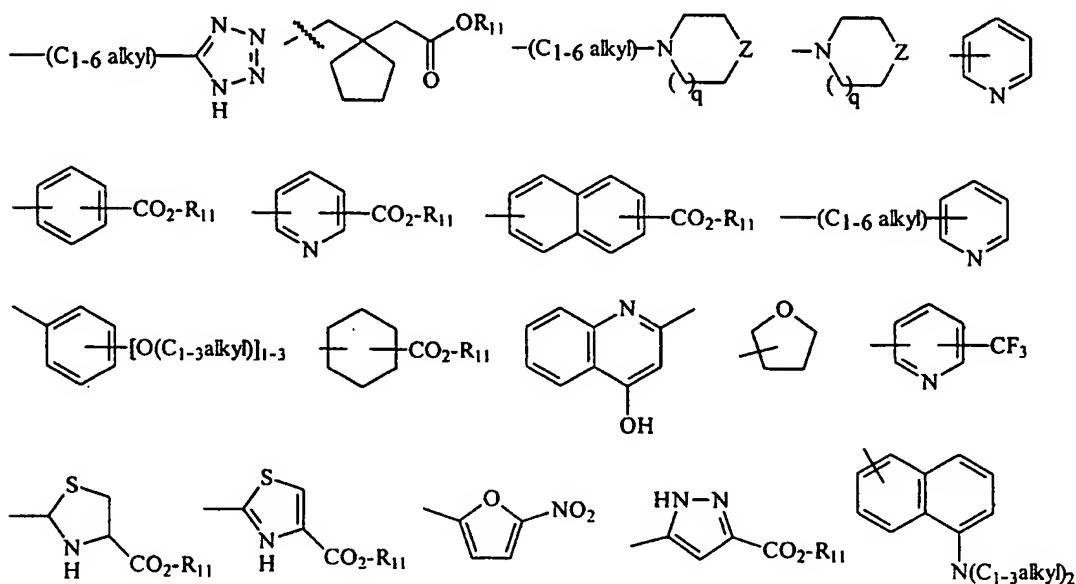
15

In yet another embodiment of the compound of formula (I), R₁ may occur one to four times and each occurrence is independently hydrogen or C₁₋₆ alkyl. Also, R₂ is hydrogen, pyridyl, C₁₋₆ alkyl, (C₁₋₆ alkyl)-CO₂-R₁₁, or -CO₂-R₁₁. Additionally, R₁ and R₂ may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:

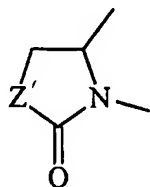
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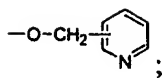
together with the carbon atom to which they are attached. In this embodiment, R_3 is hydrogen, phenyl, C_{1-6} alkyl, C_{3-6} alkenyl, C_{7-17} arylalkyl, $(C_{1-6} \text{ alkyl})-\text{CO}_2-R_{11}$, $(C_{2-6} \text{ alkenyl})-\text{CO}_2-R_{11}$, $(C_{1-6} \text{ alkyl})-\text{CO}-C_{1-6} \text{ alkyl}$, $(C_{1-6} \text{ alkyl})-\text{O}-C_{1-6} \text{ alkyl}$, $(C_{1-6} \text{ alkyl})-\text{OH}$, $(C_{1-6} \text{ alkyl})-\text{CN}$, adamantyl or one of the following:



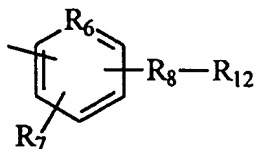
In addition, R_2 and $R_3-(Y)_6-$ may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:



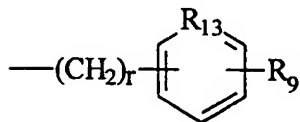
10 In this embodiment, R_4 is $-\text{O}-R_{11}$, NH_2 , NHOH , $-\text{O}-(C_{7-10} \text{ arylalkyl})$, or R_4 is of the formula:



Also, in this embodiment, R_3 is a formula of the following:



- Additionally, in this embodiment, R_6 is N or CH, R_7 is hydrogen or halogen, R_8 is -NHCO-, R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or trifluoromethyl, R_{10} is C_{1-6} alkyl, or $(C_{1-6} \text{ alkyl})\text{-OH}$, or hydrogen, R_{11} is hydrogen or C_{1-6} alkyl, R_{12} is C_{1-6} alkyl or the following formula:

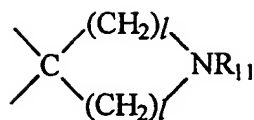


R_{13} is N or CH, W is $(C_{1-6} \text{ alkyl})$, X is S, O, or CH_2 , Y is -CO-, -C(=O)O-, -C(=O)N(R_{10})-, Z is O, CH_2 , or N- R_{11} , l is 1, 2, or 3, m is 1 or 2, n is 0 or 1, o is 0 or 1, p is 0 or 1, q is 0 or 1, and r is 0, 1, 2 or 3. In this embodiment, the following provisos apply:

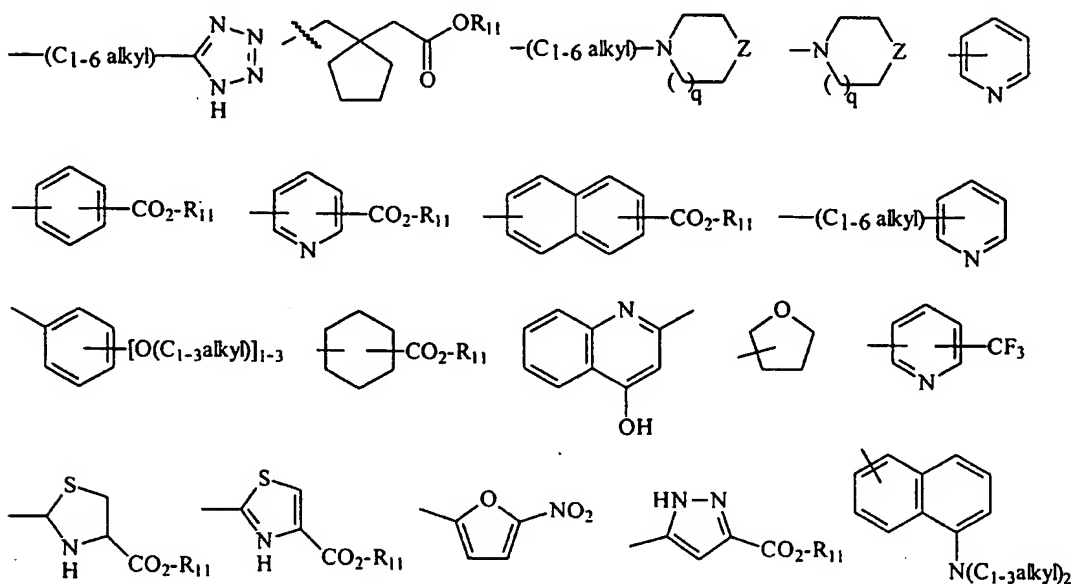
- (1) when Y is -C(=O)O-, R_3 cannot be hydrogen;
- (2) when R_4 is equal to O- $(C_4 \text{ alkyl})$, C_4 alkyl is not equal to *tert*-butyl; and
- (3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is CH_2), W is equal to CH_2 .

- In yet another embodiment of the compound of formula (I-a), R_1 may occur one to four times and each occurrence is independently hydrogen or C_{1-6} alkyl, and R_2 is hydrogen, pyridyl, C_{1-6} alkyl, $(C_{1-6} \text{ alkyl})\text{-CO}_2\text{-}R_{11}$, or $\text{-CO}_2\text{-}R_{11}$. Additionally, R_1 and R_2 may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:

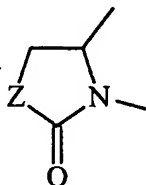
-16-



together with the carbon atom to which they are attached. Additionally, in this embodiment, R_3 is hydrogen, phenyl, C_{1-6} alkyl, C_{3-6} alkenyl, C_{7-17} arylalkyl, $(C_{1-6} \text{ alkyl})\text{-CO}_2\text{-}R_{11}$, $(C_{2-6} \text{ alkenyl})\text{-CO}_2\text{-}R_{11}$, $(C_{1-6} \text{ alkyl})\text{-CO-}C_{1-6} \text{ alkyl}$, $(C_{1-6} \text{ alkyl})\text{-O-}C_{1-6} \text{ alkyl}$, $(C_{1-6} \text{ alkyl})\text{-OH}$, $(C_{1-6} \text{ alkyl})\text{-CN}$, adamantyl or one of the following:



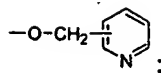
In addition, R_2 and $R_3\text{-(Y)}_o$ may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:



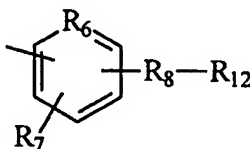
10

Additionally, R_4 is $\text{-O-}R_{11}$, NH_2 , NHOH , $\text{-O-(C}_{7-10} \text{ arylalkyl)}$, or is of the formula

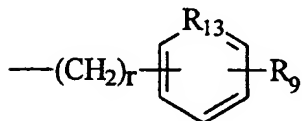
-17-



Also, in this embodiment, R_5 is a formula of the following:



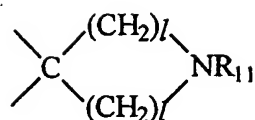
Additionally, in this embodiment, R_6 is N or CH, R_7 is hydrogen or halogen, R_8 is -NHCO-, R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or trifluoromethyl, R_{10} is C_{1-6} alkyl, or $(C_{1-6} \text{ alkyl})\text{-OH}$, or hydrogen, R_{11} is hydrogen or C_{1-6} alkyl, R_{12} is C_{1-6} alkyl or the following formula:



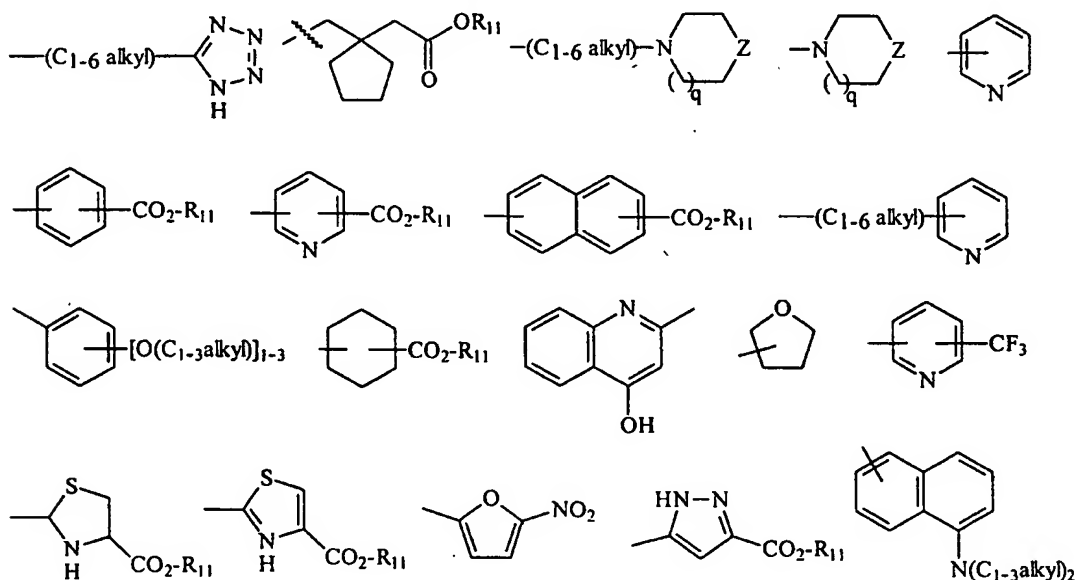
, R_{13} is N or CH, W is $(C_{1-6} \text{ alkyl})$, X is S, O, or CH_2 , Y is -CO-, -C(=O)O- or -C(=O)N(R_{10})-, Z is O, CH_2 , or N- R_{11} , l is 1, 2, or 3, m is 1 or 2, n is 0 or 1, o is 0 or 1, p is 0 or 1, q is 0 or 1, and r is 0, 1, 2 or 3. This particular embodiment of the present invention has the provisos that

- (1) when Y is -C(=O)O-, R_3 cannot be hydrogen;
- (2) when R_4 is equal to O- $(C_4 \text{ alkyl})$, $C_4 \text{ alkyl}$ is not equal to *tert*-butyl;
- (3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is CH_2), W is equal to CH_2 ; and
- (4) the compound has an IC_{50} value of less than $5\mu\text{M}$ in a Jurkat CS-1 assay and/or an IC_{50} value of less than $50\mu\text{M}$ in a Jurkat EC assay.

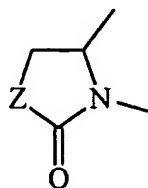
In another embodiment of the compound of formula (I), R_1 may occur one to four times and each occurrence is independently hydrogen or C_{1-6} alkyl, R_2 is hydrogen, pyridyl, C_{1-6} alkyl, $(C_{1-6} \text{ alkyl})\text{-CO}_2\text{-}R_{11}$, or $\text{-CO}_2\text{-}R_{11}$. Additionally, R_1 and R_2 may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



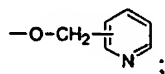
- together with the carbon atom to which they are attached. Also, in this embodiment, R_3 is hydrogen, phenyl, C_{1-6} alkyl, C_{3-6} alkenyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)- CO_2-R_{11} , $(C_{2-6}$ alkenyl)- CO_2-R_{11} , $(C_{1-6}$ alkyl)- $CO-C_{1-6}$ alkyl, $(C_{1-6}$ alkyl)- $O-C_{1-6}$ alkyl, $(C_{1-6}$ alkyl)-OH, $(C_{1-6}$ alkyl)-CN, adamantyl or one of the following:



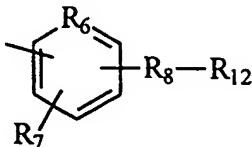
In addition, R_2 and $R_3-(Y)_o$ may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:



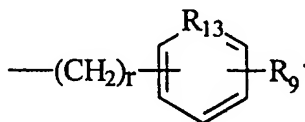
- 10 In this embodiment, R_4 is $-O-R_{11}$, NH_2 , $NHOH$, $-O-(C_{7-10}$ arylalkyl), or R_4 is of the formula:



In this embodiment, R_5 is a formula of the following:

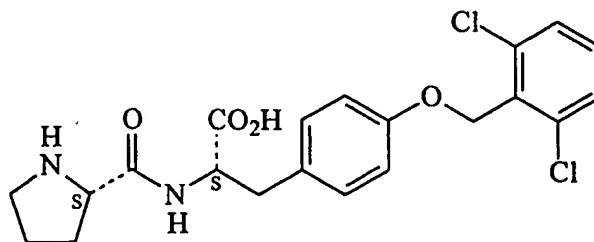


Also, in this particular embodiment, R_6 is N or CH, R_7 is hydrogen or halogen, R_8 is $-OCH_2-$, R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or trifluoromethyl, R_{10} is C_{1-6} alkyl, $(C_{1-6} \text{ alkyl})-OH$, or hydrogen, R_{11} is hydrogen or C_{1-6} alkyl, R_{12} is C_{1-6} alkyl or the following formula:

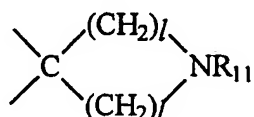


, R_{13} is N or CH, W is $(C_{1-6} \text{ alkyl})$, X is S, O, or CH_2 , Y is $-CO-$, $-C(=O)O-$ or $-C(=O)N(R_{10})-$, Z is O, CH_2 , or $N-R_{11}$, l is 1, 2, or 3, m is 1 or 2, n is 0 or 1, o is 0 or 1, p is 0 or 1, q is 0 or 1, and r is 0, 1, 2 or 3. This particular embodiment has the following provisos:

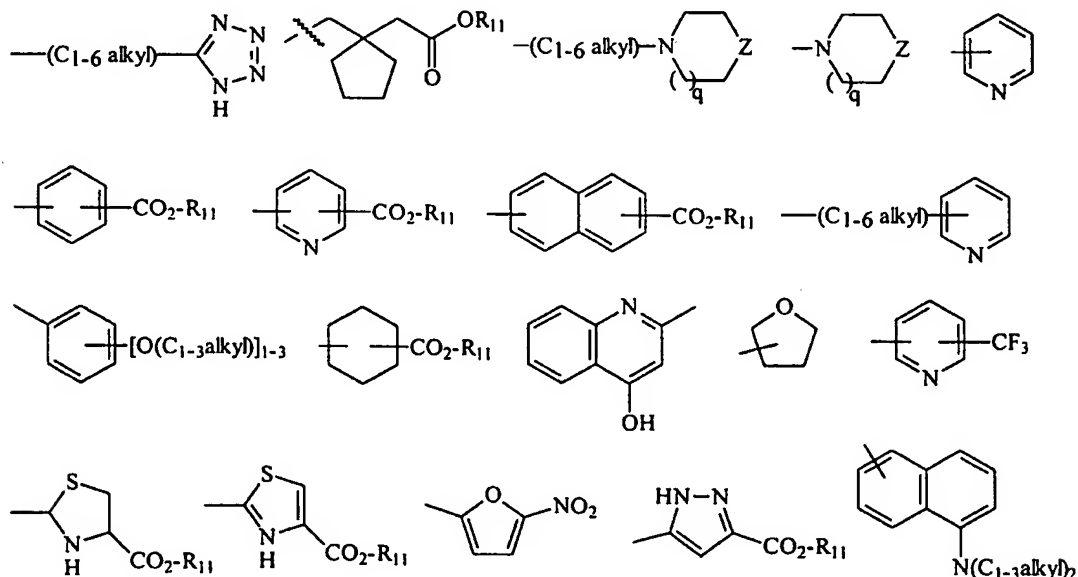
- 10 (1) when Y is $-C(=O)O-$, R_3 cannot be hydrogen;
- (2) when R_4 is equal to $O-(C_4 \text{ alkyl})$, C_4 alkyl is not equal to *tert*-butyl;
- (3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is CH_2), W is equal to CH_2 ;
- (4) when R_3 is phenyl, C_{1-6} alkyl, C_{7-17} arylalkyl, $(C_{1-6} \text{ alkyl})-CO_2-R_{11}$,
- 15 $(C_{1-6} \text{ alkyl})-O-C_{1-6} \text{ alkyl}$ or $(C_{1-6} \text{ alkyl})-OH$, o is 0; and
- (5) the compound is not



In yet another embodiment of the compound of formula (I-a), R_1 may occur one to four times and each occurrence is independently hydrogen or C_{1-6} alkyl, and R_2 is hydrogen, pyridyl, C_{1-6} alkyl, $(C_{1-6} \text{ alkyl})\text{-CO}_2\text{-R}_{11}$, or $\text{-CO}_2\text{-R}_{11}$. Additionally, R_1 and R_2 may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:

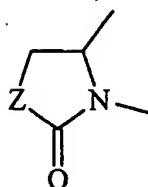


together with the carbon atom to which they are attached. Additionally, in this embodiment, R_3 is hydrogen, phenyl, C_{1-6} alkyl, C_{3-6} alkenyl, C_{7-17} arylalkyl, $(C_{1-6} \text{ alkyl})\text{-CO}_2\text{-R}_{11}$, $(C_{2-6} \text{ alkenyl})\text{-CO}_2\text{-R}_{11}$, $(C_{1-6} \text{ alkyl})\text{-CO-C}_{1-6} \text{ alkyl}$, $(C_{1-6} \text{ alkyl})\text{-O-C}_{1-6} \text{ alkyl}$, $(C_{1-6} \text{ alkyl})\text{-OH}$, $(C_{1-6} \text{ alkyl})\text{-CN}$, adamantyl or one of the following:

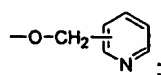


In addition, R_2 and $R_3\text{-(Y)}_0\text{-}$ may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:

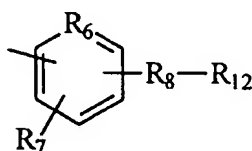
-21-



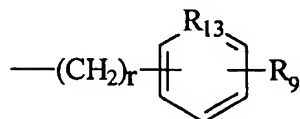
Additionally, R_4 is $-O-R_{11}$, NH_2 , $NHOH$, $-O-(C_{7-10} \text{ arylalkyl})$, or is of the formula



Also, in this embodiment, R_5 is a formula of the following:



Additionally, in this embodiment, R_6 is N or CH, R_7 is hydrogen or halogen, R_8 is $-OCH_2-$,
 5 R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or trifluoromethyl,
 R_{10} is C_{1-6} alkyl, or $(C_{1-6} \text{ alkyl})-OH$, or hydrogen, R_{11} is hydrogen or C_{1-6} alkyl, R_{12} is C_{1-6}
 alkyl or the following formula:

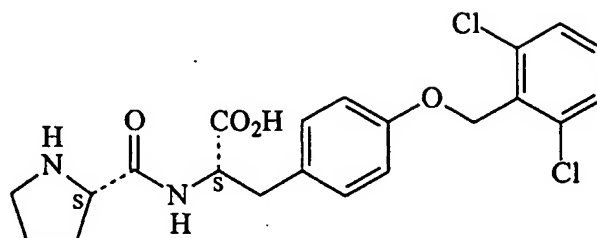


, R_{13} is N or CH, W is $(C_{1-6} \text{ alkyl})$, X is S, O, or CH_2 , Y is $-CO-$,
 $-C(=O)O-$ or $-C(=O)N(R_{10})-$, Z is O, CH_2 , or $N-R_{11}$, l is 1, 2, or 3, m is 1 or 2, n is 0 or 1, o
 10 is 0 or 1, p is 0 or 1, q is 0 or 1, and r is 0, 1, 2 or 3. This particular embodiment of the
 present invention has the provisos that:

- (1) when Y is $-C(=O)O-$, R_3 cannot be hydrogen;
- (2) when R_4 is equal to $O-(C_4 \text{ alkyl})$, C_4 alkyl is not equal to *tert*-butyl;
- (3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is
 15 CH_2), W is equal to CH_2 ;
- (4) the compound has an IC_{50} value of less than $5\mu M$ in a Jurkat CS-1 assay
 and/or an IC_{50} value of less than $50\mu M$ in a Jurkat EC assay;
- (5) when R_3 is phenyl, C_{1-6} alkyl, C_{7-17} arylalkyl, $(C_{1-6} \text{ alkyl})-CO_2-R_{11}$,

(C₁₋₆ alkyl)-O- C₁₋₆ alkyl or (C₁₋₆ alkyl)-OH, *o* is 0; and

(6) the compound is not



In the above formula (I) the absolute stereochemistry for the bond leading to R₃ is shown, however the absolute stereochemistry has not been shown for all examples which follow. It is understood that all other formulas also follow this type of absolute stereochemistry unless otherwise stated. Additionally, it is to be understood by those of skill in the art that the present invention embodies stereochemical configurations other than those shown. Specifically, the present invention embodies all configurations including the various stereoisomers. Compounds which do not meet the absolute stereochemistry in formula (I) should meet an activity threshold in various assays, to be explained below, which can ensure their efficacy as useful molecules. The present invention includes mixtures, such as racemic mixtures, which contain molecules having the claimed stereochemistry.

The desired compound of the present invention may be clinically used either in a free form or in the form of pharmaceutically acceptable salts thereof. Pharmaceutically acceptable salts include acid-addition salts with inorganic acid or organic acid (e.g., hydrochloride, sulfate, nitrate, hydrobromide, methanesulfonate, p-toluenesulfonate, acetate), salt with inorganic base, organic base or amino acid (e.g., triethylamine salt, a salt with lysine, an alkali metal salt, an alkali earth metal salt and the like).

The compound may also be formulated into a pharmaceutical composition comprising a therapeutically effective amount of the compound as defined above and a pharmaceutically acceptable carrier or diluent.

The compound can also be used for treating or preventing $\alpha_4\beta_1$ adhesion mediated conditions in a mammal such as a human. This method may comprise administering to a

mammal or a human patient an effective amount of the compound or composition as explained above.

This method can be used to treat such inflammatory conditions as rheumatoid arthritis, asthma, allergy conditions, adult respiratory distress syndrome, AIDS, cardiovascular diseases, thrombosis or harmful platelet aggregation, reocclusion following thrombolysis, allograft rejection, reperfusion injury, psoriasis, eczema, contact dermatitis and other skin inflammatory diseases, osteoporosis, osteoarthritis, atherosclerosis, neoplastic diseases including metastasis of neoplastic or cancerous growth, wound healing enhancement, treatment of certain eye diseases such as detaching retina, Type I diabetes, multiple sclerosis, systemic lupus erythematosus (SLE), inflammatory and immunoinflammatory conditions including ophthalmic inflammatory conditions and inflammatory bowel diseases, ulcerative colitis, atherosclerosis, regional enteritis and other autoimmune diseases.

As mentioned above, the compounds and compositions containing the compounds according to the present invention are particularly useful in treating or preventing $\alpha_4\beta_1$ adhesion mediated conditions in a mammal such as a human. The present inventors have found that the compounds and compositions containing the compounds according to the present invention are most useful in the treatment of asthma.

The desired compound of the present invention or pharmaceutically acceptable salts thereof may be administered either orally or parenterally, and it may be used as a suitable pharmaceutical preparation, for example, a tablet, a granule, a capsule, a powder, an injection, and an inhalation by a conventional process.

The dose of the desired compound of the present invention or a pharmaceutically acceptable salt thereof varies depending on an administration method, age, body weight, and state of a patient, but, in general, the daily dose is preferably about 0.1 to 100 mg/kg/day, however, 1 to 100 mg/kg/day may also be suitable.

Preferred routes of administration for asthma:

It is preferred that the compound of the present invention be administered in the form of an Aerosol. However, other routes of administration include intravenous, oral, intramuscular, and subcutaneous.

5 In the case of aerosol administration, compositions containing the compounds of the present invention can be prepared to provide for an excellent means for administering in aerosol form for inhalation therapy. Accordingly, the present invention will provide for self-propelling compositions containing the compounds of the present invention.

Propellants employed should be non-toxic and have a vapor pressure suitable for the
10 conditions under which administration occurs. These propellants can be fluorinated or fluorochlorinated lower saturated aliphatic hydrocarbons. The preferred propellants of this type are the halogenated alkanes containing not more than two carbon atoms and at least one fluorine atom. Illustrative of these are trichloromonofluoromethane, dichlorodifluoromethane, monochlorotrifluoromethane, dichloromonofluoromethane and
15 1,2-dichloro-1,1,2,2-tetrafluoroethane. These compounds are available from E.I. duPont de Nemours and Company under the trade name "Freon". These propellants may be employed singularly or in admixture.

In addition to the propellant, an organic solvent may also be employed. The organic solvent must be non-toxic and without undesirable effects on inhalation in the amount
20 present in the aerosol produced. In addition, the solvent should be substantially anhydrous, completely miscible with the propellant or mixture of propellants employed and have a suitable boiling point. Examples of such solvents included non-toxic aliphatic alcohols such as ethanol; ethers such as ethyl ether and vinyl ether; ketones such as acetone; and suitable halogenated lower alkanes.

25 In addition to the organic solvent, the composition may also optionally contain a non-toxic hygroscopic glycol. The glycol must be substantially miscible with the organic solvent and the propellant employed. Satisfactory glycols include propylene glycol, triethylene glycol, glycerol, butylene glycol and hexylene glycol.

The above indicated methods of administration and formulation of aerosol
30 compositions should not be viewed as limiting. The compounds of the present invention can

be formulated in anyway deemed suitable to one of ordinary skill in the art so as to obtain the desired effects.

Pharmaceutical Compositions

5 As indicated previously, the compounds of formula (I) can be formulated into pharmaceutical compositions. In determining when a compound of formula (I) is indicated for the treatment of a given disease, the particular disease in question, its severity, as well as the age, sex, weight, and condition of the subject to be treated, must be taken into consideration and this perusal is to be determined by the skill of the attendant physician.

10 For medical use, the amount of a compound of formula (I) required to achieve a therapeutic effect will, of course, vary both with the particular compound, the route of administration, the patient under treatment, and the particular disorder or disease being treated. A suitable daily dose of a compound of Formula (I), or a pharmaceutically acceptable salt thereof, for a mammalian subject suffering from, or likely to suffer from, any
15 condition as described hereinbefore is 0.1 mg to 100 mg of the compound of formula I, per kilogram body weight of the mammalian subject. In the case of systematic administration, the dose may be in the range of 0.5 to 500 mg of the compound per kilogram body weight, the most preferred dosage being 0.5 to 50 mg/kg of mammal body weight administered two to three times daily. In the case of topical administration, e.g., to the skin or eye, a suitable
20 dose may be in the range of 0.1 μ g to 100 μ g of the compound per kilogram, typically about 0.1 μ g/kg.

 In the case of oral dosing, a suitable dose of a compound of Formula (I), or a physiologically acceptable salt thereof, may be as specified in the preceding paragraph, but most preferably is from 1 mg to 10 mg of the compound per kilogram, the most preferred
25 dosage being from 1 mg to 5 mg/kg of mammal body weight, for example, from 1 to 2 mg/kg. Most preferably, a unit dosage of an orally administrable composition encompassed by the present invention contains less than about 1.0 g of a formula (I) compound.

 It is understood that formulation, both for human and veterinary use, of the present invention may be presented to the mammal by inhalation. To achieve therapeutic effect, the
30 dose may be in the range of 0.5 to 500 mg of the compound, per kg body weight. The most

preferred dosage being 0.5 to 50 mg/kg of mammal body weight administered two to three times daily.

It is understood that the ordinarily skilled physician or veterinarian will readily determine and prescribe the effective amount of a compound of formula (I) to prevent or arrest the progress of the condition for which treatment is administered. In so proceeding, the physician or veterinarian could employ relatively low doses at first, subsequently increasing the dose until a maximum response is obtained.

The compounds and compositions of the present invention can be administered to patients suffering from a condition listed herein in an amount which is effective to fully or partially alleviate undesired symptoms of the condition. The symptoms may be caused by inappropriate cell adhesion mediated by $\alpha_4\beta_1$ integrins. Such inappropriate cell adhesion would typically be expected to occur as a result of increased VCAM-1 and/or CS-1 expression on the surface of endothelial cells. Increased VCAM-1 and/or CS-1 expression can be due to a normal inflammation response or due to abnormal inflammatory states. In either case, an effective dose of a compound of the invention may reduce the increased cell adhesion due to increased VCAM-1 expression by endothelial cells. Reducing the adhesion observed in the disease state by 50% can be considered an effective reduction in adhesion. More preferably, a reduction in adhesion by 90%, is achieved. Most preferably adhesion mediated by VCAM-1/ $\alpha_4\beta_1$ and/or CS-1 interaction is abolished by an effective dose. Clinically, in some instances, effect of the compound can be observed as a decrease in white cell infiltration into tissues or a site of injury. To achieve a therapeutic effect, then, the compounds or compositions of the present invention are administered to provide a dose effective to reduce or eliminate inappropriate cell adhesion or to alleviate undesired symptoms.

While it is possible for an active ingredient to be administered alone, it is preferable to present it as a pharmaceutical formulation comprising a compound of formula (I) and a pharmaceutically acceptable carrier thereof. Such formulations constitute a further feature of the present invention.

The formulations, both for human and veterinary medical use, of the present invention comprise an active ingredient of formula (I), in association with a pharmaceutically

acceptable carrier thereof and optionally other therapeutic ingredient(s), which are generally known to be effective in treating the disease or condition encountered. The carrier(s) must be "acceptable" in the sense of being compatible with the other ingredients of the formulations and not deleterious to the recipient thereof.

5 The formulations include those in a form suitable for oral, pulmonary, ophthalmic, rectal, parenteral (including subcutaneous, intramuscular, and intravenous), intra-articular, topical, nasal inhalation (e.g., with an aerosol) or buccal administration. Such formulation are understood to include long-acting formulations known in the art.

10 The formulations may conveniently be presented in unit dosage form and may be prepared by any of the methods well known in the art of pharmacy. All methods may include the step of bringing the active ingredient into association with the carrier which constitutes one or more accessory ingredients. In general, the formulations are prepared by uniformly and intimately bringing the active ingredient into association with a liquid carrier or a finely divided solid carrier or both, and then, if necessary, shaping the product into the
15 desired form.

Formulations of the present invention suitable for oral administration may be in the form of discrete units such as capsules, cachets, tablets, or lozenges, each containing a predetermined amount of the active ingredient in the form of a powder or granules; in the form of a solution or suspension in an aqueous liquid. Formulations for other uses could
20 involve a nonaqueous liquid; in the form of an oil-in-water emulsion or a water-in-oil emulsion; in the form of an aerosol; or in the form of a cream or ointment or impregnated into a transdermal patch for use in administering the active ingredient transdermally, to a patient in need thereof. The active ingredient of the present inventive compositions may also be administered to a patient in need thereof in the form of a bolus, electuary, or paste.

25 The practitioner is referred to "Remington: The Science and Practice of Pharmacy," 19th Edition, c. 1995 by the Philadelphia College of Pharmacy and Science, as a comprehensive tome on pharmaceutical preparations.

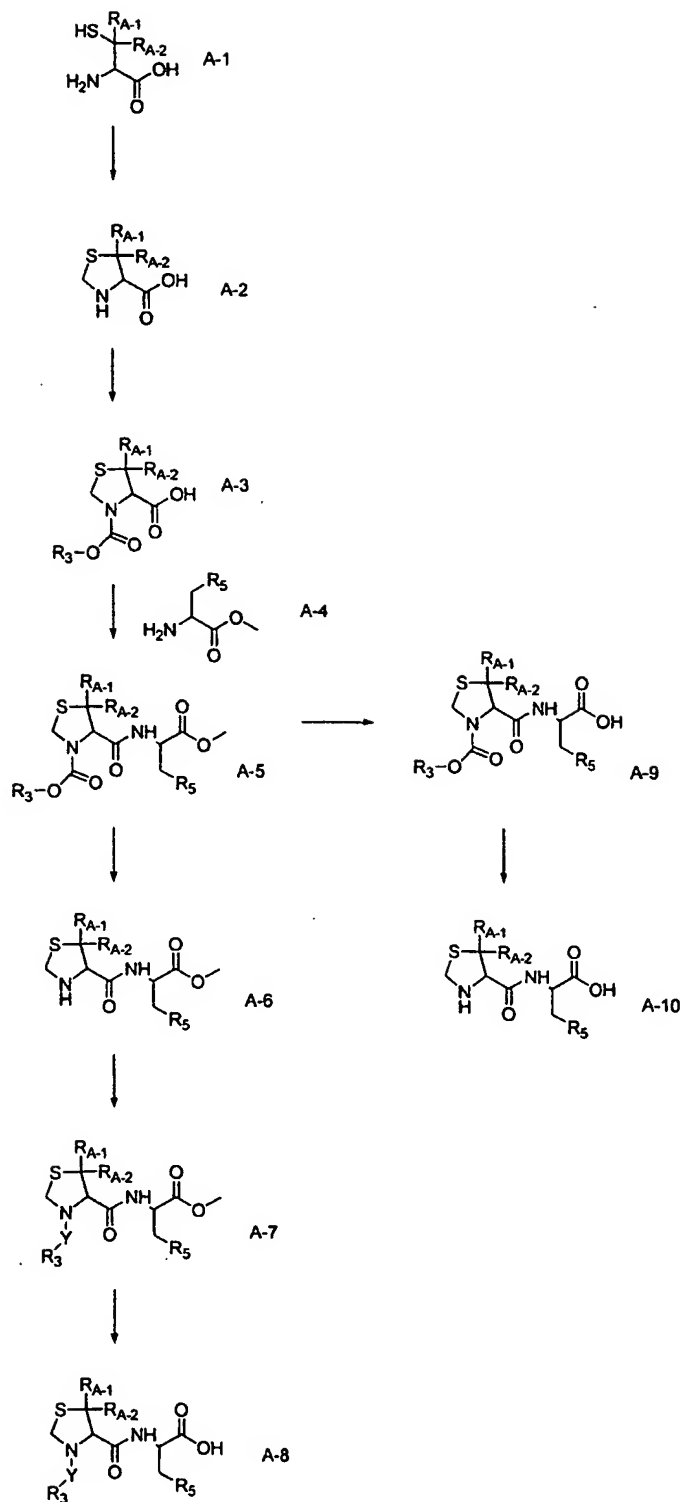
Abbreviations

	Ac ₂ O:	Acetic anhydride
	EtOAc:	Ethyl acetate
5	BCECF-AM:	2',7'-bis-(2-carboxyethyl)-5-(and 6-) carboxyfluorescein acetoxymethyl ester
	BOP-Cl:	Bis (2-oxo-3-oxazolidinyl) phosphinic chloride
	BOP Reagent:	Benzotriazol-1-yloxy-tris (dimethylamino)-phosphonium hexafluorophosphate
	DMEM:	Dulbecco's Minimal Eagle's Media
10	DMF:	Dimethyl formamide
	DIEA:	Diisopropylethylamine
	EDC:	1-(3-Dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride
	Et:	Ethyl
	EtOH:	Ethanol
15	HATU:	<i>N</i> -[(Dimethylamino)-1 <i>H</i> -1,2,3-triazolo[4,5- <i>b</i>]-pyridin-1-ylmethylene]- <i>N</i> -methylmethanaminium hexafluorophosphate <i>N</i> -oxide
	HBSS:	Hank's Balanced Salt Solution
	HBTU:	O-Benzotriazol-1-yl- <i>N,N,N',N'</i> -tetramethyluronium hexafluorophosphate
20	HOBT (HOBt):	1-Hydroxybenzotriazole
	HSA:	Human serum albumin
	LDA:	Lithium diisopropylamide
	Me:	Methyl
	meq:	milliequivalent
25	MeOH:	Methanol
	<i>n</i> -Bu:	<i>n</i> -Butyl
	NMP:	1-Methyl-2-pyrrolidinone
	PBS:	Phosphate buffered saline
	Pd-C:	Palladium on charcoal
30	Ph:	Phenyl

	SPDP:	3-(2-pyridyldithio)propionic acid N-hydroxysuccinimide ester	
	t-Bu:	t-butyl	
	THF:	Tetrahydrofuran	
	TFA:	Trifluoroacetic acid	
5	DMSO:	dimethyl sulfoxide	
	HOAt:	1-hydroxy-7-azabenzotriazole	
	DMAP:	4-dimethylaminopyridine	
	FMOC:	9-fluorenylmethoxycarbonyl	
	Bn:	benzyl	
10	PyBOP:	(benzotriazol-1-yloxy)tripyrrolidinophosphonium	hexafluoro-
		phosphate	
	BOC:	<u>tert</u> -butoxycarbonyl	

The representative compounds according to the present invention are prepared as described below. The compounds of the present invention are prepared in a similar manner.

Scheme A



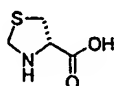
R_{A-1} and R_{A-2} are defined independently as R_1 .

Scheme A describes a general method for the preparation of examples of the formula A-5, A-6, A-7, A-8, A-9 and A-10. A commercially available or readily prepared sulfur containing amino acid of structure A-1 (for the synthesis of β,β -disubstituted cysteine amino acids see: Stanfield, G.F.; Hruby, V.J. *Synth. Commun.* **1988**, *18*, 531 and references therein) is condensed with formaldehyde to afford the thiazolidine-4-carboxylic acid of general formula A-2 (for the condensation of aldehydes with cysteine or similar sulfur containing amino acids see for example: (a) Ratner, S.; Clark, H.T. *J. Am. Chem. Soc.* **1937**, *59*, 200. (b) Lewis, N.J.; Inloes, R.L.; Hes, J.; Matthews, R.H.; Milo, G. *J. Med. Chem.* **1978**, *21*, 1070. (c) Oya, M.; Baba, T.; Kato, E.; Kawashima, Y.; Watanbe, T. *Chem. Pharm. Bull.* **1982**, *30*, 440.). Standard protection affords carbamate A-3 which is readily condensed with amino acid derivative A-4 under standard peptide synthesis conditions to provide the psuedodipeptide A-5 (for a review of procedures of peptide synthesis see: Bodansky, M.; Bodansky, A. *The Practice of Peptide Synthesis*; Springer-Verlag: Berlin, 1984). Deprotection of the carbamate from A-5 provides the useful intermediate A-6. The amine group may be reacted with a variety of electrophilic reagents such as: (1) commercially available or readily prepared sulfonyl chlorides (for the synthesis of sulfonyl chlorides see for examples: (a) Roblin, R.O.; Clapp, J.W. *J. Am. Chem. Soc.* **1950**, *72*, 4890. (b) Gilbert, E.E. in *Sulfonation and Related Reactions* Olah, G.A., Ed. John Wiley and Sons, New York; 1965. (c) Park, Y.J.; Shin, H.H.; Kin, Y.H. *Chem Lett.* **1992**, 1483. (d) Kim, D.; Ko, Y.K.; Kim, S.H. *Synthesis*, **1992**, 1203.) to afford sulfonamides of general structure A-7 where Y is SO_2 - (preparation 5); (2) carbonates or chloroformates to afford carbamates of general structure A-7 where Y is CO_2 - (preparations 2, 7, 8); (3) isocyanates to afford ureas of general structure A-7 where Y is $CONHR_3$ (preparation 9); (4) phosgene or a suitable equivalent and an amine to afford ureas of general structure A-7 where Y is $CON(C_{1-6} \text{ alkyl})R_3$ - (preparation 10, also see for example: Nowick, J.S.; Homes, D.L.; Noronha, G.; Smith, E.M.; Tram, M.N.; Huang, S. *J. Org. Chem.* **1996**, *61*, 3929.); (5) acid chlorides and carboxylic anhydrides to provide amides of structure A-7 where Y is CO - (preparation 11). Mild base hydrolysis of monoesters of general structure A-7 (preparation 6) or diesters of general structure A-7

(preparation 12) affords the acids of general structure A-8. Mild base hydrolysis of the ester of general structure A-5 provides acid A-9 (preparation 6 or 13) which may be further deprotected to afford the amino acid A-10 (preparation 14).

Preparation 1

- 5 (Scheme A, A-2: where R_{A-1} and R_{A-2} are the same and equal to H and stereochemistry is (S))

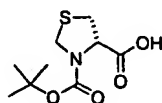


- 10 D-Cysteine hydrochloride monohydrate (A-1, where R_{A-1} and R_{A-2} are the same and equal to H and stereochemistry is (S)) (35.04 g, 0.19 mol) was dissolved in formaldehyde (40 wt% solution in water, 38 mL) and the reaction mixture allowed to stir for 18 h at ambient temperature. The mixture was cooled (0-5°C) and absolute ethanol (93 mL) and pyridine (57 mL) were added. After one hour, the precipitate was collected by filtration, washed
15 with cold absolute ethanol followed by diethyl ether and dried *in vacuo* to afford the title compound (24.6 g) as a white crystalline solid: mp 181-184°C (Lit. 194-196°C; Lewis, N.J.; Inoles, R.L.; Hes, J. *J. Med. Chem.* 1978, 21, 1070.); ^1H NMR (DMSO- d_6) δ 4.22 (1 H), 4.04 (1 H), 3.86 (1 H), 3.09 (1 H), 2.24 (1 H); MS (ESI+) for $\text{C}_4\text{H}_7\text{NO}_2\text{S}$ m/z 134.0 (M+H) $^+$.

20

Preparation 2

(Scheme A, A-3: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is *t*-butyl and stereochemistry is (S))



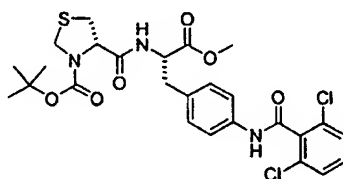
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A solution of A-2 (Scheme A where R_{A-1} and R_{A-2} are the same and equal to H and stereochemistry is (S)) (24.6 g, 0.185 mol) and di-*t*-butyl dicarbonate (44.4 g, 0.2 mol) in THF (1 L) was heated to reflux for 18 h. Volatiles were removed *in vacuo* and the residue

partitioned between ethyl acetate and 0.1 N NaOH. The aqueous layer was washed with ethyl acetate, made acidic with 1.0 N HCl (pH 3-4) and then extracted with ethyl acetate. The combined organic extracts were washed with brine, dried (Na_2SO_4), filtered and concentrated *in vacuo*. Crystallization of the white solid from hexane/methylene chloride provided the title compound (31.8 g) as white crystals: mp 132-134 °C; $[\alpha]_D^{25} = 117^\circ$ (c 0.66, ethanol); IR (mull) 3002, 1747, 1635, 1421, 1404, 1393, 1310, 1215, 1198, 1166, 1144, 1122, 894, 862, 774 cm^{-1} ; ^1H NMR (DMSO-d_6) δ 4.57 (2 H), 4.28 (1 H), 3.09 (1 H), 1.35 (9 H); ^{13}C NMR (DMSO-d_6) δ 171.8, 152.5, 79.8, 60.9, 48.4, 47.7, 33.8, 32.6, 27.7; MS (ESI+) for $\text{C}_9\text{H}_{15}\text{NO}_4\text{S}$ m/z 234.2 (M+H) $^+$; MS (ESI-) for $\text{C}_9\text{H}_{15}\text{NO}_4\text{S}$ m/z 232.1 (M-H) $^-$; Anal. Calcd for $\text{C}_9\text{H}_{15}\text{NO}_4\text{S}$: C, 46.34; H, 6.48; N, 6.00. Found: C, 46.27; H, 6.48; N, 6.03.

Preparation 3, Example 1

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester (Scheme A, A-5: where $\text{R}_{\text{A},1}$ and $\text{R}_{\text{A},2}$ are the same and equal to H, R_3 is *t*-butyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)).



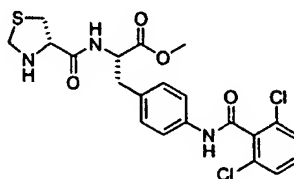
To a cooled (0-5°C) suspension of A-3 (Scheme A where $\text{R}_{\text{A},1}$ and $\text{R}_{\text{A},2}$ are the same and equal to H, R_3 is *t*-butyl and stereochemistry is (S)) (8.67 g, 37.2 mmol) and HOBT (5.69 g, 37.2 mmol) in CH_2Cl_2 (60 mL) was added a solution of EDC (7.12 g, 37.2 mmol) in CH_2Cl_2 (140 mL). After 30 min at 0-5°C, A-4 (where R_5 is 4-[(2,6-dichlorobenzoyl)-amino]phenyl, and stereochemistry is (S)) (10 g, 24.8 mmol) was added followed by 4-methylmorpholine (2.72 mL, 24.8 mmol). The reaction mixture was gradually warmed to ambient temperature, stirred an additional 18 h and diluted with CH_2Cl_2 and 0.1 N HCl. The organic layer was separated and washed with 0.1 N HCl, sat. aqueous NaHCO_3 , brine,

dried (Na_2SO_4), filtered and concentrated *in vacuo*. Flash chromatography of the residue using hexane/ethyl acetate (50%) as eluant afforded the title compound (13.9 g) as a white solid. Recrystallization from acetone/hexane afforded a crystalline solid: mp 222-224°C; IR (mull) 3282, 3254, 1738, 1714, 1707, 1678, 1662, 1610, 1562, 1545, 1431, 1414, 1287, 1256, 784 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.57 (2 H), 7.34 (5 H), 7.14 (2 H), 4.74 (3 H), 4.30 (1 H), 3.74 (3 H), 3.37 (1 H), 3.15 (3 H), 1.45 (9 H); ^{13}C NMR ($\text{DMSO}-d_6$) δ 172.1, 162.3, 153.2, 137.6, 136.9, 133.2, 131.8, 131.7, 130.1, 128.7, 119.8, 80.4, 62.2, 53.7, 52.4, 36.7, 28.3; MS (ESI+) for $\text{C}_{26}\text{H}_{29}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 604.3 ($\text{M}+\text{Na}^+$); Anal. Calcd for $\text{C}_{26}\text{H}_{29}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$: C, 53.61; H, 5.02; N, 7.21. Found C, 53.82; H, 4.81; N, 7.22.

10

Preparation 4

(Scheme A, A-6: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (*S,S*)).



15

To a cooled (5-10°C) solution of A-5 (Scheme A where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is *t*-butyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (*S,S*)) (3.3 g, 5.67 mmol) in dioxane (34 mL) was added a solution of HCl in dioxane (4 M, 140 mL) in a dropwise manner over 30 min. After an additional 30 min at 0-5°C, the ice bath was removed and the reaction mixture stirred 1 h at ambient temperature. The volatiles were removed *in vacuo* to afford the title compound (2.94 g) as a light yellow solid: ^1H NMR ($\text{DMSO}-d_6$) δ 10.70 (1 H), 9.1 (1 H), 7.52 (5 H), 7.18 (2 H), 4.64 (1 H), 4.37 (1 H), 4.21 (2 H), 3.67 (3 H), 3.10 (1 H), 2.89 (1 H), 2.70 (1 H); ^{13}C NMR ($\text{DMSO}-d_6$) δ 172.1, 162.3, 153.2, 137.6, 136.9, 133.2, 131.8, 131.7, 130.1, 128.7, 119.8, 80.4, 62.2, 53.7, 52.4, 36.7, 28.3; MS (ESI+) for $\text{C}_{21}\text{H}_{21}\text{Cl}_2\text{N}_3\text{O}_4\text{S}$ m/z 482.1 ($\text{M}+\text{H}^+$).

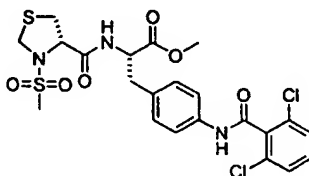
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Preparation 5 and Example 2.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-(methylsulfonyl)-4-thiazolidinyl]carbonyl]-L-phenylalanine methyl ester

(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is methyl, Y is SO_2 ,

5 R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



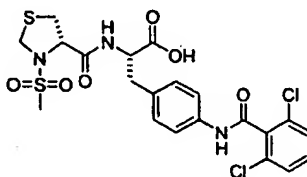
To a cooled (0-5°C) solution of A-6 (Scheme A, where R_{A-1} and R_{A-2} are the same and
 10 equal to H, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)) (2.0 g, 3.86 mmol) in anhydrous THF (50 mL) was added methanesulfonyl chloride (2.99 mL, 38.6 mmol) followed by pyridine (6.24 mL, 77.2 mmol). After 1 h at 0-5°C, the ice bath was removed and the solution stirred at ambient temperature for 2 h then diluted with ethyl acetate and 0.25 N HCl. The layers were separated and the organic layer washed with sat.
 15 aqueous $NaHCO_3$, brine, dried ($MgSO_4$), filtered and concentrated *in vacuo*. Purification of the residue by flash chromatography using ethyl acetate/methylene chloride/hexane (1:1:1) and isopropanol (0.1%) as eluant afforded the title compound (1.99 g) as an amorphous powder: IR (mull) 1742, 1666, 1605, 1562, 1534, 1515, 1432, 1413, 1344, 1327, 1269, 1218, 1195, 1156, 780 cm^{-1} ; 1H NMR (300 MHz, $CDCl_3$) δ 7.55 (3 H), 7.33 (3 H), 7.12 (3 H), 4.84 (1 H), 4.69 (1 H), 4.61 (1 H), 4.29 (1 H), 3.74 (3 H), 3.50 (1 H),
 20 3.29 (1 H), 3.14 (2 H), 2.92 (3 H); ^{13}C NMR (75 MHz, $CDCl_3$) δ 171.2, 168.1, 162.4, 136.5, 135.9, 132.3, 132.2, 130.8, 130.0, 128.1, 120.6, 64.9, 53.2, 52.5, 51.8, 37.2, 34.1; MS (ESI+) for $C_{22}H_{23}Cl_2N_3O_6S_2$ m/z 559.8 (M+H) $^+$; HRMS (FAB) calcd for $C_{22}H_{23}Cl_2N_3O_6S_2+H_1$ 560.0483, found 560.0504; Anal. Calcd for $C_{22}H_{23}Cl_2N_3O_6S_2$: C, 47.15; H, 4.14; N, 7.50. Found: C, 46.88; H, 4.32; N, 7.16.
 25

Preparation 6 and Example 3.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-(methylsulfonyl)-4-thiazolidinyl]carbonyl]-L-phenylalanine

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is methyl, Y is SO_2 ,

5 R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



To a cooled (0-5°C) solution of A-7 (Scheme A, where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is methyl, Y is SO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)) (1.75 g, 3.12 mmol) in THF (100 mL) and water (10 mL) was added an 0.1 N aqueous solution of NaOH (34.3 mL, 3.43 mmol) via a syringe pump over 1 h. After an additional 45 min at 0-5°C, the reaction mixture was diluted with ethyl acetate and acidified with 0.25 N HCl to a pH of ca. 3. The organic layer was separated, washed with water and concentrated *in vacuo*. Purification of the residue by flash chromatography using methylene chloride and methanol (0-5%) as eluant provided a solid which was lyophilized from glacial acetic acid to provide the title compound (1.42g) as an amorphous powder: $[\alpha]_D^{25} = 103^\circ$ (c 0.97, ethanol); IR (mull) 3291, 1736, 1666, 1605, 1562, 1534, 1516, 1432, 1414, 1339, 1270, 1195, 1154, 799, 780 cm^{-1} . 1H NMR (300 MHz, CD_3OD) δ 7.58 (2 H), 7.45 (3 H), 7.25 (2 H), 4.72 (2 H), 4.37 (1 H), 3.17 (5 H), 2.99 (3 H); ^{13}C NMR (75 MHz, CD_3OD) δ 174.1, 171.6, 165.2, 138.3, 137.7, 134.8, 133.4, 132.4, 131.2, 129.4, 121.6, 66.2, 55.0, 52.9, 37.9, 37.5, 35.7; MS (ESI+) for $C_{21}H_{21}Cl_2N_3O_6S_2$ m/z 545.8 (M+H) $^+$; HRMS (FAB) calcd for $C_{21}H_{21}Cl_2N_3O_6S_2+H_1$ 546.0327, found 546.0358. Anal. Calcd for $C_{21}H_{21}Cl_2N_3O_6S_2$: C, 46.16; H, 3.87; N, 7.69.

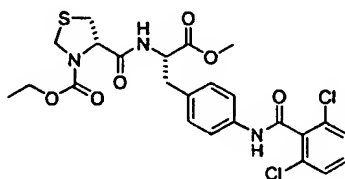
25 Found: C, 46.24; H, 4.04; N, 7.33.

Preparation 7 and Example 4.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester

(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is ethyl, Y is CO₂-,

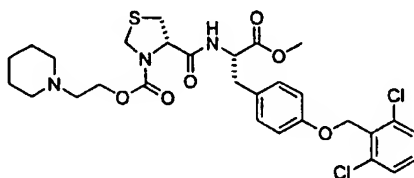
5 R₃ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



To a cooled (0-5°C) solution of A-6 (where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)) (1.25 g, 2.40 mmol) in anhydrous THF (30 mL) was added ethyl chloroformate (340 µL, 3.60 mmol) followed by triethylamine (810 µL, 5.79 mmol). After 1 h at 0-5°C, the ice bath was removed and the solution stirred at ambient temperature for 2 h then diluted with ethyl acetate and 0.25 N HCl. The layers were separated and the organic layer washed with sat. aqueous NaHCO₃, brine, dried (MgSO₄), filtered and concentrated *in vacuo*. Purification of the residue by flash chromatography using ethyl acetate/methylene chloride/hexane (1:1:1) and isopropanol (0.1%) as eluant afforded the title compound (1.10 g) as an amorphous powder: ¹H NMR (300 MHz, CDCl₃) δ 7.59 (2 H), 7.30 (3 H), 7.10 (2 H), 4.81 (1 H), 4.72 (2 H), 4.38 (1 H), 4.11 (2 H), 3.19 (4 H), 1.25 (3 H); ¹³C NMR (75 MHz, CDCl₃) δ 171.6, 163.0, 136.9, 136.2, 132.3, 132.1, 130.7, 129.8, 128.0, 120.3, 63.0, 62.8, 57.1, 53.2, 52.5, 37.0, 14.4; MS (ESI+) for C₂₄H₂₃Cl₂N₃O₆S *m/z* 554.2 (M+H)⁺; MS (FAB) *m/z* (rel. intensity) 554 (MH⁺, 99), 557 (29), 556 (76), 555 (45), 554 (99), 349 (35), 245 (27), 175 (35), 173 (52), 160 (93), 88 (38); HRMS (FAB) calcd for C₂₄H₂₃Cl₂N₃O₆S + H₁, 554.0919, found 554.0908. Anal. Calcd for C₂₄H₂₃Cl₂N₃O₆S: C, 51.99; H, 4.55; N, 7.58. Found: C, 52.05; H, 4.67; N, 7.44.

Preparation 8 and Example 5 .

[S-(R*,R*)]-4-[[[1-[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(1-piperidiny)ethyl] ester (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is 2-(1-piperidiny)ethyl, Y is CO₂-, R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).



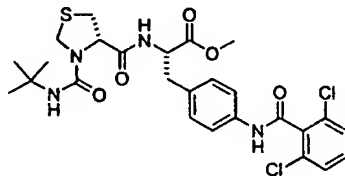
The title compound was prepared by a modification of the literature procedure of Ghosh, A.K.; Duong, T.T.; McKee, S.P.; Thompson, W.J. *Tetrahedron Lett.* **1992**, *33*, 2781. To a solution of 1-(2-hydroxyethyl)piperidine (5.11 g, 39.6 mmol) in CH₃CN (220 mL) at ambient temperature was added N,N-disuccinimidyl carbonate (10.13 g, 39.6 mmol) and triethylamine (16.6 mL, 118.8 mmol). The solution was stirred at room temperature for 4 h and concentrated *in vacuo* to give a viscous oil. The oil was dissolved in a minimal amount of methylene chloride (50 mL) and added to a solution of A-6 (Scheme A, where R_{A-1} and R_{A-2} are the same and equal to H, R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)) (2.0 g, 3.96 mmol), triethylamine (0.60 mL) and DMAP (5 mg) in CH₂Cl₂ (10 mL). The solution was stirred overnight and an additional 5 equivalents of carbonate in methylene chloride (10 mL) [prepared as described above from N,N-disuccinimidyl carbonate (5.6 g, 19.8 mmol), triethylamine (8.3 mL, 59.4 mmol), and 1-(2-hydroxyethyl)piperidine (2.56 g, 19.8 mmol)] were added. After 3 h at room temperature, propylamine (30 mL, 0.71 mol) was slowly added (*exothermic*) and the solution diluted with CH₂Cl₂. The resulting solution was stirred vigorously for 15 min. and diluted with water. The organic layer was separated and washed with 0.1 M HCl, sat. aqueous NaHCO₃, dried (MgSO₄), filtered and concentrated *in vacuo*. Purification of the residue by flash chromatography using ethyl acetate/methylene chloride (3:1) as eluant

followed by trituration in hexanes afforded the title compound (1.54 g, 62%) as a white powder: IR (mull) 1745, 1704, 1660, 1553, 1512, 1435, 1426, 1397, 1303, 1245, 1227, 1212, 1173, 1019, 765 cm^{-1} ; ^1H NMR (300 MHz, DMSO-d_6) δ 8.42 (1 H), 7.54 (2 H), 7.44 (1 H), 7.13 (2 H), 6.94 (2 H), 5.17 (2 H), 4.59 (2 H), 4.48 (1 H), 4.26 (1 H), 4.02 (2 H), 3.63 (3 H), 3.19 (2 H), 3.19 (1 H), 2.82 (3 H), 2.35 (4 H), 1.36 (6 H); ^{13}C NMR (75 MHz, DMSO-d_6) δ 172.2, 170.0, 157.7, 153.9, 136.5, 132.2, 132.0, 130.8, 130.1, 129.2, 114.8, 65.3, 63.6, 57.3, 54.6, 53.9, 52.4, 36.4, 26.0, 24.3; MS (ESI+) for $\text{C}_{29}\text{H}_{35}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 623.9 ($\text{M}+\text{H}$) $^+$; Anal. Calcd for $\text{C}_{29}\text{H}_{35}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$: C, 55.77; H, 5.65; N, 6.73. Found: C, 55.48; H, 5.73; N, 6.91.

Preparation 9 and Example 6.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-[(1,1-dimethylethyl)amino]carbonyl]-4-thiazolidinyl]carbonyl]-L-phenylalanine methyl ester

(Scheme A, A-7: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is *t*-butyl, Y is CONH-, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*, *S*)).

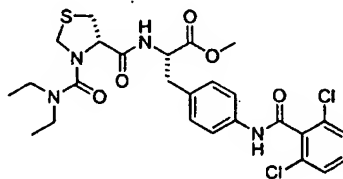


To a cooled (0-5°C) solution of A-6 (Scheme A, where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*, *S*)) (140 mg, 0.27 mmol) in anhydrous THF (6 mL) was added *tert*-butyl isocyanate (0.62 mL, 5.4 mmol) followed by 4-dimethylaminopyridine (5 mg, 0.04 mmol). After 0.5 h at 0-5°C, the ice bath was removed and the solution stirred at ambient temperature for 16 h. Additional *tert*-butyl isocyanate (0.62 mL, 5.4 mmol) was added and the solution warmed to 50°C for 4 h. The reaction mixture was cooled to room temperature and diluted with ethyl acetate and 0.25 N HCl. The layers separated and the organic layer washed with sat. aqueous NaHCO_3 , brine, dried (MgSO_4), filtered and concentrated *in vacuo*. Purification of the residue by flash chromatography using ethyl acetate/methylene chloride (1:1:1) and

isopropanol (0.1%) as eluant afforded the title compound (150 mg) as an amorphous powder: ^1H NMR (300 MHz, CDCl_3) δ 7.93 (1 H), 7.54 (2 H), 7.26 (4 H), 7.09 (2 H), 4.74 (2 H), 4.66 (1 H), 4.41 (1 H), 4.23 (1 H), 3.70 (3 H), 3.28 (1 H), 3.09 (3 H), 1.31 (9 H); ^{13}C NMR (75 MHz, CDCl_3) δ 171.4, 170.6, 162.4, 155.5, 136.3, 135.9, 132.5, 132.3, 130.8, 129.9, 128.1, 120.6, 62.7, 53.2, 52.4, 51.5, 49.0, 37.0, 32.9, 29.2; MS (ESI+) for $\text{C}_{26}\text{H}_{30}\text{Cl}_2\text{N}_4\text{O}_5\text{S}$ m/z 581.0 (M+H) $^+$, 603.0 (M+Na) $^+$; MS (FAB) m/z (rel. intensity) 581 (MH $^+$, 23), 482 (50), 97 (36), 88 (36), 83 (45), 69 (99), 57 (81), 55 (79), 43 (50), 43 (69), 41 (50). HRMS (FAB) calcd for $\text{C}_{26}\text{H}_{30}\text{Cl}_2\text{N}_4\text{O}_5\text{S} + \text{H}^+$ 581.1392, found 581.1376.

Preparation 10 and Example 7.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-[(diethylamino)carbonyl]-4-thiazolidinyl]carbonyl]-L-phenylalanine methyl ester
(Scheme A, A-7: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is ethyl, Y is $\text{CON}(\text{CH}_2\text{CH}_3)_2$, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).

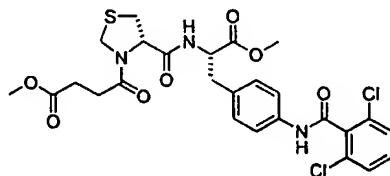


The title compound was prepared by a modification of the literature procedure of Majer, P.; Randad, R.S. *J. Org. Chem.* 1994, 59, 1937. A cooled (0-5°C) solution of A-6
(Scheme A, where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)) (200 mg, 0.39 mmol) and triethylamine (56 μL , 0.40 mmol) in anhydrous methylene chloride (10 mL) was added to triphosgene (47 mg, 0.16 mmol) followed by additional triethylamine (56 μL , 0.40 mmol). After 0.5 h at 0-5°C, the ice bath was removed and the solution stirred at ambient
temperature for 2 h. The solution was re-cooled (0-5°C) and diethylamine (1.20 mL, 11.70 mmol) and 4-dimethylaminopyridine (5 mg, 0.04 mmol) were added. After 0.5 h at 0-5°C, the ice bath was removed and the solution stirred at ambient temperature for 16 h.

The solution was concentrated *in vacuo* and the residue partitioned between ethyl acetate and 0.25 N HCl. The layers were separated and the organic layer washed with sat. aqueous NaHCO₃, brine, dried (MgSO₄), filtered and concentrated *in vacuo*. Purification of the residue by flash chromatography using ethyl acetate/methylene chloride/hexane (1:1:1) and isopropanol (0.1%) as eluant afforded the title compound (200 mg) as an amorphous solid: ¹H NMR (300 MHz, CDCl₃) δ 7.86 (1 H), 7.55 (2 H), 7.30 (3 H), 7.03 (2 H), 6.90 (1 H), 5.09 (1 H), 4.83 (1 H), 4.33 (2 H), 3.76 (3 H), 3.34 (3 H), 3.11 (5 H), 1.08 (6 H); ¹³C NMR (CDCl₃) δ 171.1, 169.6, 162.3, 162.2, 136.5, 135.9, 132.3, 132.1, 130.9, 129.9, 128.1, 120.3, 64.7, 53.4, 52.7, 52.5, 42.0, 37.0, 32.4, 13.1; MS (ESI+) for C₂₆H₃₀N₄O₅S *m/z* 580.9 (M+H)⁺; HRMS (EI) calcd for C₂₆H₃₀Cl₂N₄O₅S 580.1314, found 580.1297. Anal. Calcd for C₂₆H₃₀Cl₂N₄O₅S: C, 53.70; H, 5.20; N, 9.63. Found: C, 53.63; H, 5.33; N, 9.36.

Preparation 11 and Example 8.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-γ-oxo-3-thiazolidinebutanoic acid 3-methyl ester (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is CH₂CH₂CO₂CH₃, Y is CO, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (*S,S*)).



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To a cooled (0-5°C) solution of A-6 (Scheme A, where R_{A-1} and R_{A-2} are the same and equal to H, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S,S*)) (1.03 g, 1.72 mmol) in anhydrous CH₂Cl₂ (25 mL) was added triethylamine (460 μL, 3.27 mmol) followed by methyl succinyl chloride (320 μL, 2.58 mmol). After 1 h at 0-5°C, the ice bath was removed and the solution stirred at ambient temperature for 2 h then diluted with 1 N HCl. The organic layer was separated, washed with sat. aqueous NaHCO₃, brine, dried (Na₂SO₄), filtered and concentrated *in vacuo*. Crystallization of the

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yellow solid from ethanol/water provided the title compound (824 mg) as a light yellow solid: mp 221-223 °C; IR (mull) 3275, 1748, 1731, 1687, 1626, 1610, 1561, 1542, 1517, 1430, 1416, 1326, 1268, 1224, 1193 cm^{-1} ; ^1H NMR ($\text{DMSO}-d_6$) δ 10.64 (1 H), 8.59 (1 H), 8.24 (1 H), 7.50 (5 H), 7.16 (2 H), 4.75 (2 H), 4.51 (2 H), 4.23 (1 H), 3.63 (3 H), 3.56 (3 H), 2.87 (5 H); ^{13}C NMR ($\text{DMSO}-d_6$) δ 172.9, 171.6, 169.9, 169.5, 161.9, 137.2, 136.5, 133.0, 132.8, 131.3, 129.8, 128.3, 119.4, 61.6, 53.6, 52.1, 51.4, 48.6, 36.4, 35.7, 35.2, 33.1, 29.0, 28.9, 28.5; MS (ESI+) for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 596.0 ($\text{M}+\text{H}$) $^+$; MS (ESI-) for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 593.9 ($\text{M}-\text{H}$) $^-$; Anal. Calcd for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$: C, 52.35; H, 4.56; N, 7.04. Found: C, 52.11; H, 4.47; N, 6.96.

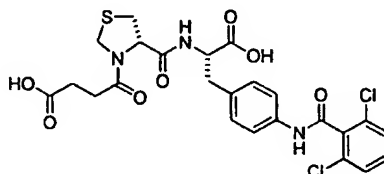
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Preparation 12 and Example 9.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)-amino]phenyl]ethyl]amino]carbonyl]- γ -oxo-3-thiazolidinebutanoic acid

(Scheme A, A-8: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is $\text{CH}_2\text{CH}_2\text{CO}_2\text{CO}_2\text{H}$, Y is CO, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)).

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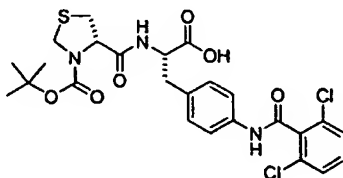
To a cooled (0-5°C) solution of A-7 (Scheme A where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is $\text{CH}_2\text{CH}_2\text{CO}_2\text{CH}_3$, Y is CO, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)) (130 mg, 0.22 mmol) in anhydrous THF (5 mL) and MeOH (1 mL) was added an aqueous (2 mL) solution of lithium hydroxide monohydrate (23 mg, 0.55 mmol) via a syringe pump over 1 h. After an additional 1 h at 0-5°C, the ice bath was removed and the solution stirred 2 h at ambient temperature. The reaction mixture was diluted with ethyl acetate and 0.1 N HCl and the organic layer was separated, washed with water, dried (Na_2SO_4), filtered and concentrated *in vacuo*. Lyophilization of the residue from glacial acetic acid afforded the title compound (101 mg) as a white amorphous

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powder: ^1H NMR (CD_3CN) δ 8.87 (1 H), 7.55 (2 H), 7.42 (3 H), 7.22 (2 H), 7.16 (1 H), 4.95 (1 H), 4.52 (3 H), 3.12 (5 H), 2.55 (5 H); ^{13}C NMR (CD_3CN) δ 173.8, 171.7, 171.1, 169.8, 162.5, 136.7, 136.0, 133.5, 131.7, 131.3, 130.1, 129.2, 128.2, 119.7, 62.4, 53.6, 48.8, 36.2, 32.4, 29.1, 28.6; MS (ESI-) for $\text{C}_{24}\text{H}_{23}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 566.1 (M-H) $^-$; Anal. Calcd for $\text{C}_{24}\text{H}_{23}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$: C, 50.31; H, 4.13; N, 7.33. Found: C, 50.13; H, 4.37; N, 6.93.

Preparation 13 and Example 10.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]-phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester (Scheme A, A-9: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is *t*-butyl, Y is CO_2^- , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)).



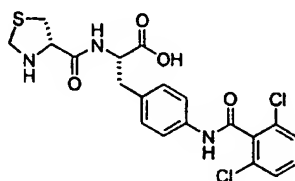
To a cooled (0-5°C) of A-5 (Scheme A where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is *t*-butyl, Y is CO_2^- , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)) (150 mg, 0.26 mmol) in anhydrous THF (5 mL) and MeOH (1 mL) was added an aqueous (2 mL) solution of lithium hydroxide monohydrate (14 mg, 0.325 mmol) via syringe pump over 1 h. After an additional 1 h at 0-5°C, the ice bath was removed and the solution stirred 2 h at ambient temperature. The reaction mixture was diluted with ethyl acetate and 0.1 N HCl, the organic layer separated and washed with water, dried (Na_2SO_4), filtered and concentrated *in vacuo*. Lyophilization of the residue from glacial acetic acid afforded the title compound (142 mg) as an amorphous powder: IR (mull) 3285, 1735, 1666, 1606, 1562, 1539, 1516, 1432, 1413, 1394, 1326, 1259, 1219, 1195, 1161 cm^{-1} ; ^1H NMR (DMF-d_7) δ 10.71 (1 H), 8.36 (1 H), 7.91 (2 H), 7.72 (3 H), 7.47 (2 H), 4.88 (3 H), 4.51 (1 H), 3.40 (3 H), 3.22 (2 H), 1.57 (9 H); ^{13}C NMR (DMF-d_7) δ 173.3, 163.1, 162.9, 162.7, 162.3, 154.0, 138.3, 137.5, 134.2, 132.3, 131.9, 130.6, 128.9, 120.0, 80.9, 63.1, 54.5, 49.9, 37.4, 28.3; MS (FAB) m/z (rel. intensity) 568 (MH $^+$, 23),

570 (14), 568 (23), 471 (13), 470 (65), 469 (23), 468 (99), 466 (23), 175 (19), 88 (17), 57 (42); HRMS (FAB) calcd for $C_{25}H_{27}Cl_2N_3O_6S+H_1$ 568.1075, found 568.1071; MS (ESI-) for $C_{25}H_{27}Cl_2N_3O_6S$ m/z 565.8 (M-H); Anal. Calcd for $C_{25}H_{27}Cl_2N_3O_6S \cdot 0.26 H_2O$: C, 52.38; H, 4.84; N, 7.33. Found: C, 52.07; H, 5.12; N, 7.46; % Water (KF): 0.83.

Preparation 14 and Example 11.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-4-thiazolidinyl]carbonyl]-L-phenylalanine monohydrochloride salt

(Scheme A, A-10: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)).

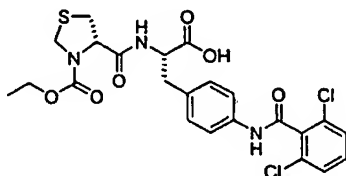


To a cooled (5-10°C) solution of A-9 (Scheme A where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is *t*-butyl, Y is CO_2^- , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)) (193 mg, 0.34 mmol) in dioxane (2 mL) was added a solution of HCl in dioxane (4 M, 8 mL) in a dropwise manner over 30 min. After an additional 3.5 h at 0-5°C, the reaction mixture was concentrated *in vacuo*. Lyophilization of the residue from water afforded the title compound (158 mg) as an amorphous powder: IR (mull) 3248, 3191, 3048, 1731, 1664, 1605, 1577, 1562, 1541, 1516, 1431, 1414, 1327, 1195, 799 cm^{-1} ; 1H NMR (DMF- d_7) δ 9.11 (1 H), 7.76 (2 H), 7.60 (3 H), 7.35 (2 H), 4.66 (2 H), 4.46 (2 H), 3.55 (3 H), 3.24 (2 H), 3.10 (2 H); ^{13}C NMR (DMF- d_7) δ 172.8, 167.6, 163.1, 162.9, 162.7, 162.3, 138.5, 137.5, 133.8, 132.3, 131.9, 130.6, 128.9, 120.0, 63.8, 54.8, 50.0, 37.3; MS (FAB) m/z (rel. intensity) 468 (MH $^+$, 99), 544 (18), 528 (15), 472 (13), 471 (16), 470 (70), 469 (24), 468 (99), 175 (14), 173 (16), 88 (18); HRMS (FAB) calcd for $C_{20}H_{19}Cl_2N_3O_4S+H_1$ 468.0551, found 468.0556.

Example 12.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester

- 5 (Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is ethyl, Y is CO₂, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



- 10 Example 12 was prepared from example 4 by the procedure described in preparation 6. Physical properties as follows: mp 118-121°C; $[\alpha]_D^{25} = 106^\circ$ (c 0.88, ethanol); IR (mull) 3283, 3196, 1665, 1606, 1561, 1539, 1516, 1431, 1414, 1345, 1327, 1271, 1219, 1195, 799 cm⁻¹; ¹H NMR (300 MHz, CD₃OD) δ 7.59 (2 H), 7.44 (3 H), 7.22 (2 H), 4.69 (1 H), 4.64 (1 H), 4.41 (1 H), 3.24 (3 H), 2.95 (2 H), 1.26 (3 H); ¹³C NMR (75 MHz, CDCl₃) δ 172.4, 171.5, 163.0, 154.8, 136.7, 136.6, 132.4, 132.1, 130.6, 129.9, 127.9, 120.3, 63.0, 62.7, 53.1, 36.7, 14.3; MS (FAB) *m/z* (rel. intensity) 540 (MH⁺, 59), 544 (12), 543 (17), 542 (53), 540 (59), 160 (32), 123 (15), 118 (20), 107 (99), 95 (11), 23 (21); HRMS (FAB) calcd for C₂₃H₂₃Cl₂N₃O₆S + H⁺, 540.0762, found 540.0730. Anal. Calcd for C₂₃H₂₃Cl₂N₃O₆S: C, 51.12; H, 4.29; N, 7.78. Found: C, 50.77; H, 4.43; N, 7.68.

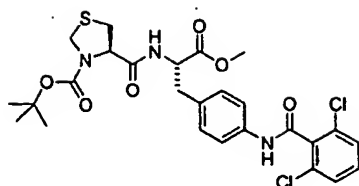
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Example 13.

[R-(R*,S*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is *t*-butyl, Y is CO₂, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (R, S)).

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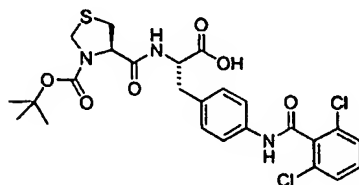


Example 13 was prepared as described in Scheme A from L-cysteine using di-*t*-butyl dicarbonate to form the requisite carbamate. Physical data as follows: IR (mull) 1746,
 5 1666, 1606, 1562, 1538, 1516, 1432, 1413, 1324, 1267, 1260, 1216, 1195, 1162, 799 cm^{-1} ;
 ^1H NMR (CDCl_3) δ 7.56 (2 H), 7.46 (1 H), 7.33 (3 H), 7.13 (2 H), 6.94 (1 H), 4.75 (3 H),
 4.25 (1 H), 3.75 (3 H), 3.39 (1 H), 3.14 (3 H), 1.43 (9 H); ^{13}C NMR ($\text{DMSO}-d_6$) δ 171.6,
 171.5, 170.7, 170.1, 161.7, 152.6, 137.0, 136.9, 136.2, 132.6, 132.6, 131.2, 131.0, 129.3,
 128.1, 119.1, 79.7, 78.2, 61.5, 53.4, 53.3, 51.8, 49.3, 49.1, 35.8, 27.6; MS (ESI+) for
 10 $\text{C}_{26}\text{H}_{29}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 604 ($\text{M}+\text{Na}$) $^+$; MS (ESI-) for $\text{C}_{26}\text{H}_{29}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 580 ($\text{M}-\text{H}$) $^-$; Anal.
 Calcd for $\text{C}_{26}\text{H}_{29}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.17 \text{H}_2\text{O}$: C, 53.34; H, 5.05; N, 7.18. Found: C, 53.47; H,
 5.14; N, 7.15. % Water (KF): 0.51.

Example 14.

[R-(R^* , S^*)]-4-[[[1-Carboxy-2-[4-[(2,6-
 15 dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-
 (1,1-dimethylethyl) ester

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is *t*-butyl, Y is CO_2 ,
 R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (R , S)).



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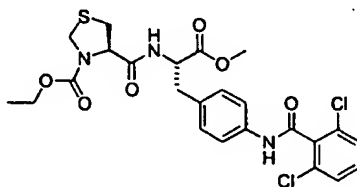
Example 14 was prepared from example 13 by the procedure described in preparation 6.
 Physical data as follows: IR (mull) 3285, 1665, 1607, 1562, 1538, 1516, 1432, 1413, 1394,
 1327, 1259, 1217, 1195, 1162, 799 cm^{-1} ; ^1H NMR ($\text{DMSO}-d_6$) δ 12.70 (1 H), 10.67 (1 H),

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8.14 (1 H), 7.51 (5 H), 7.20 (2 H), 4.56 (1 H), 4.35 (3 H), 2.98 (3 H), 1.22 (9 H); ^{13}C NMR (DMSO- d_6) δ 172.5, 170.1, 169.9, 161.6, 152.6, 136.9, 136.3, 133.0, 131.2, 131.0, 129.4, 128.1, 119.1, 79.8, 79.7, 61.6, 53.4, 49.2, 48.3, 35.9, 27.6, 20.9; HRMS (FAB) calcd for $\text{C}_{25}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_6\text{S}+\text{H}^+$ 568.1075, found 568.1058; MS (ESI+) for $\text{C}_{25}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 567.8 (M+H) $^+$; MS (ESI-) for $\text{C}_{25}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 565.8 (M-H) $^-$; Anal. Calcd for $\text{C}_{25}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.24 \text{ H}_2\text{O}$: C, 52.43; H, 4.84; N, 7.34. Found: C, 52.23; H, 4.76; N, 7.24. % Water (KF): 0.75.

Example 15.

[R-(R*,S*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester
(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is ethyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (R, S)).

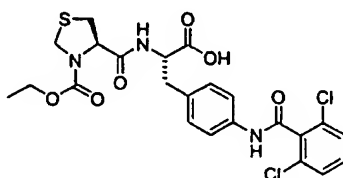


Example 15 was prepared as described in Scheme A from L-cysteine using ethyl chloroformate to form the requisite carbamate. Physical data as follows: IR (mull) 1744, 1666, 1606, 1561, 1538, 1515, 1445, 1431, 1414, 1345, 1325, 1270, 1216, 1194, 1184 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.55 (2 H), 7.36 (4 H), 7.13 (2 H), 6.95 (1 H), 4.74 (3 H), 4.21 (3 H), 3.75 (3 H), 3.40 (1 H), 3.13 (3 H), 1.26 (3 H); ^{13}C NMR (CDCl_3) δ 171.2, 169.4, 162.3, 136.2, 135.8, 132.4, 132.2, 131.0, 130.3, 130.1, 129.9, 128.2, 128.0, 127.9, 120.2, 62.9, 62.7, 53.2, 52.5, 37.1, 14.5, 14.3; MS (ESI+) for $\text{C}_{24}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 553.8 (M+H) $^+$; MS (ESI-) for $\text{C}_{24}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 551.8 (M-H) $^-$; Anal. Calcd for $\text{C}_{24}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.24 \text{ H}_2\text{O}$: C, 51.59; H, 4.60; N, 7.52. Found: C, 51.89; H, 4.62; N, 7.51. % Water (KF): 0.77.

Example 16.

[R-(R*,S*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester

- 5 (Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is ethyl, Y is CO₂, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (R, S)).

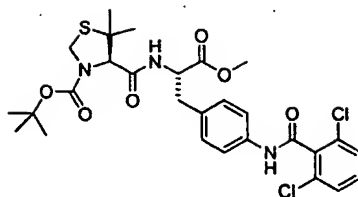


- 10 Example 16 was prepared from example 15 by the procedure described in preparation 6. Physical data as follows: IR (mull) 3287, 1664, 1606, 1561, 1539, 1516, 1445, 1431, 1414, 1346, 1327, 1271, 1217, 1195, 799 cm⁻¹; ¹H NMR (DMSO-d₆) δ 12.80 (1 H), 10.69 (1 H), 8.26 (1 H), 7.51 (5 H), 7.19 (2 H), 4.62 (2 H), 4.37 (2 H), 3.94 (2 H), 2.96 (3 H), 1.11 (3 H); ¹³C NMR (DMSO-d₆) δ 172.5, 169.6, 161.7, 153.5, 136.9, 136.3, 133.0, 131.2, 131.0, 129.5, 128.1, 119.1, 61.2, 53.4, 52.5, 35.9, 22.3, 14.3; MS (ESI+) for C₂₃H₂₃Cl₂N₃O₆S m/z 540.0 (M+H)⁺; MS (ESI-) for C₂₃H₂₃Cl₂N₃O₆S m/z 538.0 (M-H)⁻; HRMS (FAB) calcd for C₂₃H₂₃Cl₂N₃O₆S+H₁ 540.0762, found 540.0775; Anal. Calcd for C₂₃H₂₃Cl₂N₃O₆S • 0.34 H₂O: C, 50.54; H, 4.37; N, 7.69. Found: C, 50.53; H, 4.48; N, 7.59. % Water (KF): 1.13.

- 20 Example 17.

[R-(R*,S*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-5,5-dimethyl-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester

- (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to CH₃, R₃ is *t*-butyl, Y is CO₂, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (R, S)).

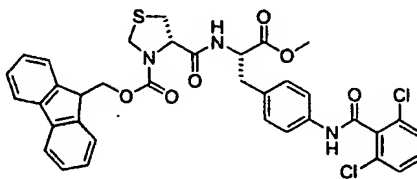


Example 17 was prepared as described in Scheme A from L-penicillamine using di-t-butyl dicarbonate to form the requisite carbamate. Physical data as follows: IR (mull) 1747,
 5 1666, 1606, 1562, 1537, 1516, 1432, 1413, 1324, 1268, 1259, 1213, 1195, 1161, 1142 cm^{-1} ;
 ^1H NMR (CDCl_3) δ 7.56 (2 H), 7.34 (4 H), 7.21 (2 H), 6.44 (1 H), 4.94 (1 H), 4.60 (2 H),
 4.08 (1 H), 3.70 (3 H), 3.10 (2 H), 1.53 (3 H), 1.42 (9 H), 1.25 (3 H); ^{13}C NMR (CDCl_3) δ
 171.4, 162.3, 136.3, 135.8, 132.6, 132.4, 131.0, 130.2, 130.0, 128.2, 120.5, 120.3, 120.2,
 72.7, 63.9, 60.4, 52.9, 52.3, 48.3, 38.0, 30.3, 28.1, 23.9, 21.0, 14.2; MS (ESI+) for
 10 $\text{C}_{28}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 630.7 ($\text{M}+\text{Na}$) $^+$; Anal. Calcd for $\text{C}_{28}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.13 \text{H}_2\text{O}$: C,
 54.87; H, 5.47; N, 6.86. Found: C, 54.54; H, 5.55; N, 6.54. % Water (KF): 0.38.

Example 18.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[(9H-fluoren-1-yl)methyl]
 15 ester

(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 9-fluorenylmethyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



20

Example 18 was prepared as described in Scheme A from D-cysteine using 9-fluorenylmethyl chloroformate to form the requisite carbamate. Physical data as follows:
 IR (mull) 3280, 1750, 1692, 1671, 1604, 1560, 1538, 1515, 1441, 1430, 1422, 1346,
 1320, 1222, 1118 cm^{-1} ; ^1H NMR ($\text{DMSO}-d_6$) δ 8.59 (1 H), 7.87 (2 H), 7.49 (12 H), 4.65

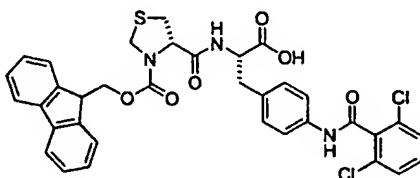
(3 H), 4.26 (4 H), 3.52 (3 H), 2.96 (3 H); ^{13}C NMR (DMSO- d_6) δ 171.4, 161.8, 143.5, 140.6, 137.0, 136.3, 132.5, 131.2, 131.1, 129.5, 128.1, 127.6, 127.1, 125.2, 125.1, 120.0, 119.2, 70.6, 70.0, 63.8, 63.2, 53.3, 53.1, 46.4, 36.2, 25.4; MS (ESI+) for $\text{C}_{36}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 703.9 (M+H) $^+$; Anal. Calcd for $\text{C}_{36}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.1 \text{ H}_2\text{O}$: C, 61.23; H, 4.45; N, 5.95.

5 Found: C, 61.18; H, 4.56; N, 5.89. % Water (KF): 0.22.

Example 19.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[(9H-fluoren-1-yl)methyl] ester

10 (Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 9-fluorenylmethyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).PNU-



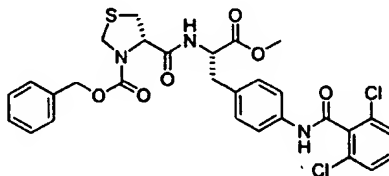
15 Example 19 was prepared from example 18 by the procedure described in preparation 6. Physical data as follows: IR (mull) 1672, 1606, 1561, 1533, 1517, 1431, 1413, 1347, 1324, 1269, 1218, 1195, 1116, 760, 742 cm^{-1} ; ^1H NMR (DMSO- d_6) δ 10.71 (1 H), 8.32 (1 H), 7.87 (2 H), 7.47 (13 H), 7.16 (2 H), 4.62 (2 H), 4.15 (5 H), 2.90 (4 H); ^{13}C NMR (DMSO- d_6) δ 172.4, 169.2, 161.8, 143.6, 140.6, 137.0, 136.3, 133.0, 131.3, 131.1, 129.7, 129.6, 128.9, 128.2, 127.7, 127.1, 126.8, 125.2, 121.3, 120.1, 120.0, 119.2, 53.4, 48.4, 46.4, 36.4, 29.5, 20.0; MS (ESI+) for $\text{C}_{35}\text{H}_{29}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 690.1 (M+H) $^+$; Anal. Calcd for $\text{C}_{35}\text{H}_{29}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.4 \text{ H}_2\text{O}$: C, 60.25; H, 4.30; N, 6.02. Found: C, 59.88; H, 4.47; N, 5.75. % Water (KF): 1.02.

Example 20.

25 [S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-phenylmethyl ester

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(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is phenylmethyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



5

Example 20 was prepared as described in Scheme A from D-cysteine using benzyl chloroformate to form the requisite carbamate. Physical data as follows: IR (mull) 1748, 1694, 1690, 1673, 1610, 1561, 1542, 1517, 1441, 1430, 1408, 1355, 1324, 1269, 1217 cm^{-1} ; 1H NMR ($CDCl_3$) δ 7.50 (2 H), 7.33 (9 H), 7.09 (2 H), 6.75 (1 H), 5.19 (2 H), 4.78 (3 H), 4.38 (3 H), 3.73 (3 H), 3.20 (3 H); ^{13}C NMR ($DMSO-d_6$) δ 171.4, 162.3, 136.3, 135.9, 135.7, 132.5, 132.4, 131.0, 130.0, 128.7, 128.4, 128.2, 128.1, 120.4, 68.2, 63.3, 53.2, 52.5, 37.2; MS (ESI+) for $C_{29}H_{27}Cl_2N_3O_6S$ m/z 637.8 ($M+Na$) $^+$; MS (ESI-) for $C_{29}H_{27}Cl_2N_3O_6S$ m/z 613.8 ($M-H$) $^-$; Anal. Calcd for $C_{29}H_{27}Cl_2N_3O_6S \cdot 0.1 H_2O$: C, 56.39; H, 4.43; N, 6.80. Found: C, 56.31; H, 4.67; N, 6.71. % Water (KF): 0.19.

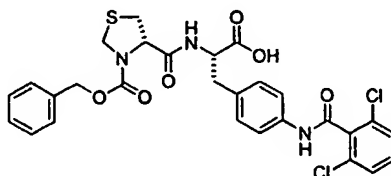
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Example 21.

[S-(R^* , R^*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-phenylmethyl ester

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is phenylmethyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).

20

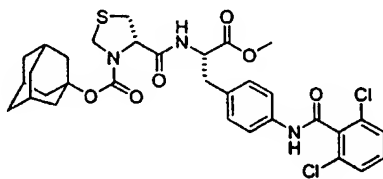


Example 21 was prepared from example 20 by the procedure described in preparation 6. Physical data as follows: IR (mull) 3290, 3034, 1666, 1606, 1562, 1537, 1516, 1431, 1413, 1351, 1326, 1270, 1215, 1195, 799 cm^{-1} ; ^1H NMR ($\text{DMSO}-d_6$) δ 12.85 (1 H), 10.65 (1 H), 7.39 (10 H), 7.18 (2 H), 4.98 (2 H), 4.65 (2 H), 4.55 (1 H), 4.33 (1 H), 3.06 (1 H), 2.83 (2 H); ^{13}C NMR (CD_3OD) δ 172.6, 163.7, 154.4, 136.8, 136.2, 136.1, 131.9, 130.9, 129.6, 128.1, 127.9, 127.8, 127.7, 127.6, 120.0, 67.5, 66.7, 53.4, 36.5; MS (ESI-) for $\text{C}_{28}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 599.7 (M-H); MS (FAB) m/z (rel. intensity) 602 (MH+, 99), 678 (37), 604 (74), 603 (33), 602 (99), 560 (32), 558 (48), 468 (35), 466 (51), 371 (50), 91 (73); HRMS (FAB) calcd for $\text{C}_{28}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_6\text{S}+\text{H}$, 602.0919, found 602.0913; Anal. Calcd for $\text{C}_{28}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.23 \text{H}_2\text{O}$: C, 55.45; H, 4.23; N, 6.93. Found: C, 55.53; H, 4.46; N, 6.88. % Water (KF): 0.67.

Example 22.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(tricyclo[3.3.1.1^{3,7}]dec-1-yl) ester

(Scheme A, A-7: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is 1-adamantyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



20

Example 22 was prepared as described in Scheme A from D-cysteine using 1-adamantyl fluoroformate to form the requisite carbamate. Physical data as follows: IR (mull) 3284, 3271, 1747, 1690, 1684, 1666, 1557, 1532, 1436, 1412, 1355, 1298, 1194, 1053, 799 cm^{-1} ; ^1H NMR ($\text{DMSO}-d_6$) δ 10.67 (1 H), 8.41 (1 H), 7.53 (5 H), 7.17 (2 H), 4.51 (3 H), 4.23 (1 H), 3.63 (3 H), 3.22 (1 H), 3.04 (1 H), 2.90 (1 H), 2.75 (1 H), 2.01 (9 H), 1.56 (6 H); ^{13}C NMR ($\text{DMSO}-d_6$) δ 172.1, 162.3, 152.7, 137.6, 136.9, 133.2, 131.8, 131.6, 130.0, 128.7, 119.8, 53.8, 52.4, 36.7, 36.1, 30.6; MS (ESI+) for $\text{C}_{32}\text{H}_{35}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 659.7

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(M+H)⁺; MS (ESI-) for C₃₂H₃₅Cl₂N₃O₆S *m/z* 657.7 (M-H)⁻; MS (FAB) *m/z* (rel. intensity) 660 (MH⁺, 8), 662 (5), 660 (8), 618 (6), 616 (8), 480 (5), 173 (7), 136 (11), 135 (99), 123 (14), 93 (8); Anal. Calcd for C₃₂H₃₅Cl₂N₃O₆S • 0.04 H₂O: C, 58.12; H, 5.35; N, 6.35. Found: C, 58.19; H, 5.62; N, 6.25. % Water (KF): 0.10.

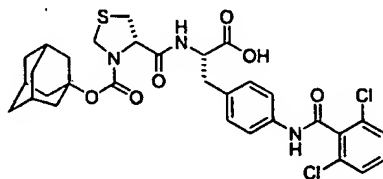
5

Example 23.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(tricyclo[3.3.1.1^{3,7}]dec-1-yl) ester

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is 1-adamantyl, Y is CO₂, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).

10



Example 23 was prepared from example 22 by the procedure described in preparation 6.

Physical data as follows: IR (mull) 3287, 1667, 1606, 1562, 1537, 1516, 1431, 1412, 1353, 1326, 1299, 1274, 1220, 1194, 1049 cm⁻¹; ¹H NMR (DMSO-d₆) δ 12.70 (1 H), 10.67 (1 H), 8.21 (1 H), 7.58 (5 H), 7.17 (2 H), 4.51 (3 H), 4.22 (1 H), 3.20 (1 H), 3.04 (1 H), 2.88 (1 H), 2.76 (1 H), 2.07 (9 H), 1.12 (6 H); ¹³C NMR (DMSO-d₆) δ 173.1, 162.3, 152.7, 137.5, 136.9, 133.8, 131.8, 131.6, 130.1, 128.7, 119.7, 80.0, 53.8, 36.8, 36.1, 30.6; MS (ESI+) for C₃₁H₃₃Cl₂N₃O₆S *m/z* 645.8 (M+H)⁺; HRMS (FAB) calcd for C₃₁H₃₃Cl₂N₃O₆S+H₁ 646.1545, found 646.1564; Anal. Calcd for C₃₁H₃₃Cl₂N₃O₆S • 0.29 H₂O: C, 57.13; H, 5.19; N, 6.45. Found: C, 56.82; H, 5.21; N, 6.32. % Water (KF): 0.80.

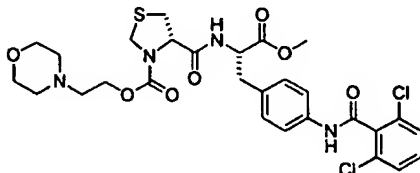
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Example 24.

[S-(R*,R*)]-4-[[[1-[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(4-morpholinyl)ethyl] ester

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(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 2-(4-morpholinyl)ethyl, Y is CO_2^- , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*, *S*)).



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Example 24 was prepared as described in Scheme A from D-cysteine using 4-(2-hydroxyethyl)morpholine to form the requisite carbamate. Physical data as follows: IR (mull) 1745, 1705, 1679, 1605, 1536, 1515, 1431, 1414, 1344, 1323, 1269, 1216, 1194,
 10 1183, 1117 cm^{-1} ^1H NMR (300 MHz, CDCl_3) δ 7.82 (1 H), 7.56 (2 H), 7.31 (3 H), 7.12 (2 H), 6.95 (1 H), 4.65 (3 H), 4.39 (3 H), 3.74 (3 H), 3.69 (3 H), 3.34 (1 H), 3.18 (3 H), 2.763 (5 H); ^{13}C NMR (75 MHz, CDCl_3) δ 171.5, 169.8, 162.9, 154.4, 136.8, 136.1, 132.1, 130.6, 129.7, 127.9, 120.2, 66.3, 62.7, 57.0, 53.4, 53.2, 52.4, 36.7, 29.5; MS (ESI-) for $\text{C}_{28}\text{H}_{32}\text{Cl}_2\text{N}_4\text{O}_7\text{S}$ m/z 636.8 (M-H) $^-$; HRMS (FAB) calcd for $\text{C}_{28}\text{H}_{32}\text{Cl}_2\text{N}_4\text{O}_7\text{S} + \text{H}_1$
 15 639.1447, found 639.1419. Anal. Calcd for $\text{C}_{28}\text{H}_{32}\text{Cl}_2\text{N}_4\text{O}_7\text{S}$: C, 52.58; H, 5.04; N, 8.76. Found: C, 52.47; H, 5.17; N, 8.69.

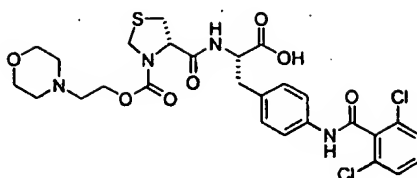
Example 25.

[S-(R^* , R^*)]4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(4-morpholinyl)ethyl] ester

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(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 2-(4-morpholinyl)ethyl, Y is CO_2^- , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*, *S*)).

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Example 25 was prepared from example 24 by the procedure described in preparation 6.

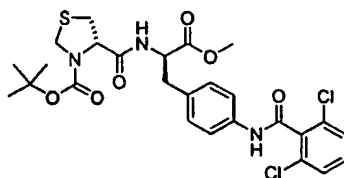
Physical data as follows: IR (mull) 3278, 1667, 1606, 1562, 1541, 1515, 1431, 1413,

- 5 1351, 1326, 1270, 1195, 1134, 1118, 799 cm^{-1} ; ^1H NMR (300 MHz, DMSO-d_6) δ 10.64 (1 H), 8.25 (1 H), 7.50 (5 H), 7.16 (2 H), 4.60 (2 H), 4.44 (1 H), 4.27 (1 H), 4.06 (2 H), 3.51 (4 H), 3.43 (2 H), 3.29 (4 H), 2.42 (4 H); ^{13}C NMR (75 MHz, DMSO-d_6) δ 173.0, 172.4, 169.8, 162.3, 137.5, 136.8, 133.5, 131.7, 130.1, 128.6, 119.7, 66.6, 63.3, 57.0, 53.7, 36.7, 21.5; MS (FAB) m/z (rel. intensity) 625 (MH^+ , 55), 629 (9), 628 (14), 627 (39), 626 (21), 625 (55), 308 (7), 141 (19), 114 (99), 113 (24), 100 (7); HRMS (FAB) calcd for $\text{C}_{27}\text{H}_{30}\text{Cl}_2\text{N}_4\text{O}_7\text{S} + \text{H}^+$ 625.1290, found 625.1309.
- 10

Example 26.

[S-(R*,S*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester

- 15 (Scheme A, A-7: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is *t*-butyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, R)).



- 20 Example 26 was prepared as described in Scheme A from D-cysteine using di-*t*-butyl dicarbonate to form the requisite carbamate. Physical data as follows: IR (mull) 3293, 1746, 1666, 1606, 1562, 1538, 1516, 1432, 1413, 1324, 1260, 1216, 1195, 1162, 799 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.55 (2 H), 7.37 (4 H), 7.14 (2 H), 4.89 (1 H), 4.66 (2 H), 4.25 (1 H), 3.75 (3 H), 3.39 (1 H), 3.24 (3 H), 1.44 (9 H); ^{13}C NMR (CDCl_3) δ 171.2, 169.8, 162.3,

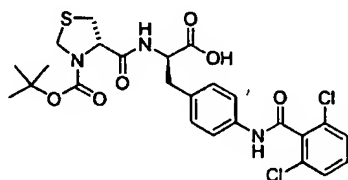
-56-

136.3, 135.8, 132.4, 132.3, 131.0, 130.2, 130.1, 129.9, 128.2, 128.1, 127.9, 120.5, 120.2, 120.1, 82.2, 53.2, 52.5, 37.3, 31.0, 28.4, 28.2; MS (ESI+) for $C_{26}H_{29}Cl_2N_3O_6S$ m/z 603.9 ($M+Na$)⁺; MS (ESI-) for $C_{26}H_{29}Cl_2N_3O_6S$ m/z 580.0 ($M-H$)⁻; HRMS (FAB) calcd for $C_{26}H_{29}Cl_2N_3O_6S+H$, 582.1232, found 582.1231. Anal. Calcd for $C_{26}H_{29}Cl_2N_3O_6S \cdot 0.26$ H₂O: C, 53.18; H, 5.07; N, 7.16. Found: C, 52.78; H, 5.14; N, 6.91. % Water (KF): 0.66.

Example 27.

[S-(R*,S*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is *t*-butyl, Y is CO₂, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*, *R*)).

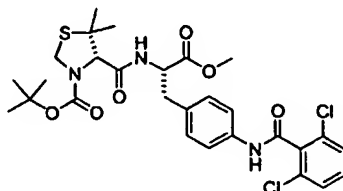


Example 27 was prepared from example 26 by the procedure described in preparation 6. Physical data as follows: IR (mull) 3286, 1665, 1606, 1562, 1538, 1516, 1432, 1413, 1394, 1326, 1259, 1216, 1195, 1161, 799 cm⁻¹; ¹H NMR (DMSO-d₆) δ 12.70 (1 H), 10.63 (1 H), 8.13 (1 H), 7.51 (5 H), 7.20 (2 H), 4.45 (4 H), 2.96 (3 H), 1.23 (9 H); ¹³C NMR (DMSO-d₆) δ 173.1, 170.5, 162.3, 153.2, 137.5, 136.9, 131.8, 131.6, 130.0, 128.7, 119.7, 80.4, 62.2, 54.0, 49.8, 36.5, 28.3, 21.5; MS (ESI+) for $C_{25}H_{27}Cl_2N_3O_6S$ m/z 567.9 ($M+H$)⁺; MS (ESI-) for $C_{25}H_{27}Cl_2N_3O_6S$ m/z 565.9 ($M-H$)⁻; HRMS (EI) calcd for $C_{25}H_{27}Cl_2N_3O_6S$ 567.0997, found 568.1096. Anal. Calcd for $C_{25}H_{27}Cl_2N_3O_6S \cdot 0.34$ H₂O: C, 52.82; H, 4.79; N, 7.39. Found: C, 52.17; H, 4.90; N, 7.25. % Water (KF): 1.07.

Example 28.

[S-(R*,R*)]-4-[[[1-[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-5,5-dimethyl-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester

(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to CH_3 , R_3 is *t*-butyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*, *S*)).



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Example 28 was prepared as described in Scheme A from D-penicillamine using di-*t*-butyl dicarbonate to form the requisite carbamate. Physical data as follows: IR (mull) 1744, 1707, 1688, 1678, 1657, 1606, 1562, 1541, 1516, 1431, 1414, 1326, 1253, 1161, 1140 cm^{-1} ; 1H NMR ($CDCl_3$) δ 7.58 (2 H), 7.36 (3 H), 7.14 (2 H), 6.55 (1 H), 4.87 (1 H), 4.56 (2 H), 4.10 (1 H), 3.72 (3 H), 3.08 (2 H), 1.53 (3 H), 1.44 (9 H), 1.40 (3 H); ^{13}C NMR ($CDCl_3$) δ 171.6, 162.3, 153.6, 136.4, 135.8, 132.6, 132.4, 131.0, 130.1, 129.9, 129.8, 128.2, 120.7, 120.5, 120.3, 81.7, 73.0, 52.8, 52.6, 52.4, 48.4, 39.8, 39.5, 37.6, 30.3, 28.3, 28.0, 23.5; MS (ESI-) for $C_{28}H_{33}Cl_2N_3O_6S$ m/z 607.9 (M-H) $^-$; MS (FAB) m/z (rel. intensity) 610 (MH $^+$, 6), 512 (26), 510 (44), 117 (30), 115 (16), 99 (16), 87 (16), 59 (99), 57 (27), 57 (20), 41 (23); HRMS (FAB) calcd for $C_{28}H_{33}Cl_2N_3O_6S+H$, 610.1545, found 610.1501; Anal. Calcd for $C_{28}H_{33}Cl_2N_3O_6S \cdot 0.07 H_2O$: C, 54.97; H, 5.46; N, 6.87. Found: C, 54.92; H, 5.54; N, 7.11. % Water (KF): 0.21.

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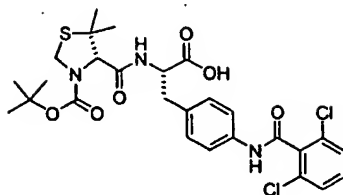
Example 29.

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[S-(R^* , R^*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-5,5-dimethyl-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to CH_3 , R_3 is *t*-butyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*, *S*)).

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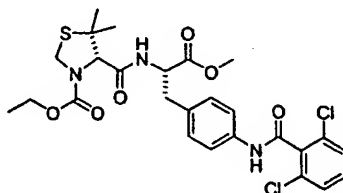
Example 29 was prepared from example 28 by the procedure described in preparation 6.

Physical data as follows: IR (mull) 1739, 1666, 1606, 1562, 1535, 1516, 1432, 1413,

- 5 1394, 1325, 1270, 1260, 1194, 1160, 799 cm^{-1} ; ^1H NMR (DMSO-d_6) δ 12.62 (1 H), 10.62 (1 H), 8.18 (1 H), 7.51 (5 H), 7.19 (2 H), 4.50 (3 H), 4.24 (1 H), 2.91 (2 H), 1.33 (12 H), 1.04 (3 H); ^{13}C NMR (DMSO-d_6) δ 172.8, 168.7, 161.9, 153.2, 137.2, 136.5, 133.1, 131.4, 131.3, 129.6, 128.3, 119.5, 80.0, 70.9, 53.7, 48.5, 37.0, 30.7, 28.1, 27.9, 24.6; MS (ESI+) for $\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 595.9 ($\text{M}+\text{H}^+$); MS (ESI-) for $\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 593.8 ($\text{M}-\text{H}$);
- 10 MS (FAB) m/z (rel. intensity) 596 (MH^+ , 19), 672 (17), 596 (19), 499 (15), 498 (60), 497 (26), 496 (99), 494 (35), 173 (20), 116 (27), 57 (48); Anal. Calcd for $\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.27 \text{H}_2\text{O}$: C, 53.93; H, 5.29; N, 6.99. Found: C, 53.73; H, 5.39; N, 7.10. % Water (KF): 0.80.

Example 30.

- 15 [S-(R^*, R^*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-5,5-dimethyl-3-thiazolidinecarboxylic acid 3-ethyl ester (Scheme A, A-7: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to CH_3 , R_3 is ethyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



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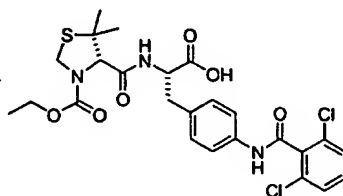
Example 30 was prepared as described in Scheme A from D-penicillamine using ethyl chloroformate to form the requisite carbamate. Physical data as follows: IR (mull) 3292, 1748, 1666, 1606, 1562, 1538, 1516, 1445, 1431, 1414, 1341, 1325, 1271, 1212, 1194 cm^{-1}

¹H NMR (CDCl₃) δ 7.57 (2 H), 7.37 (4 H), 7.11 (2 H), 6.49 (1 H), 4.86 (1 H), 4.59 (2 H), 4.14 (3 H), 3.75 (3 H), 3.09 (2 H), 1.60 (3 H), 1.54 (3 H), 1.23 (3 H); ¹³C NMR (CDCl₃) δ 171.6, 162.3, 154.5, 136.3, 135.8, 132.6, 132.4, 131.1, 130.2, 130.1, 129.8, 128.2, 120.5, 120.4, 72.7, 62.5, 52.7, 52.5, 37.3, 30.2, 23.7, 14.6, 14.1; MS (ESI+) for C₂₆H₂₉Cl₂N₃O₆S *m/z* 581.9 (M+H)⁺; MS (ESI-) for C₂₆H₂₉Cl₂N₃O₆S *m/z* 579.8 (M-H)⁻; HRMS (EI) calcd for C₂₆H₂₉Cl₂N₃O₆S 581.1154, found 581.1132; Anal. Calcd for C₂₆H₂₉Cl₂N₃O₆S • 0.16 H₂O: C, 53.35; H, 5.05; N, 7.18. Found: C, 53.74; H, 5.12; N, 7.12. % Water (KF): 0.49.

Example 31.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-5,5-dimethyl-3-thiazolidinecarboxylic acid 3-ethyl ester

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to CH₃, R₃ is ethyl, Y is CO₂, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



Example 31 was prepared from example 30 by the procedure described in preparation 6.

Physical data as follows: IR (mull) 3287, 3070, 1666, 1606, 1562, 1538, 1516, 1431, 1414, 1342, 1328, 1271, 1213, 1194, 799 cm⁻¹; ¹H NMR (DMSO-d₆) δ 12.58 (1 H), 10.63 (1 H), 8.22 (1 H), 7.51 (5 H), 7.20 (2 H), 4.51 (3 H), 3.95 (3 H), 3.04 (1 H), 2.86 (1 H), 1.35 (3 H), 1.16 (6 H); ¹³C NMR (DMSO-d₆) δ 223.3, 184.1, 183.9, 172.8, 168.4, 161.7, 136.9, 136.3, 133.1, 131.2, 131.0, 129.4, 128.1, 119.2, 70.6, 61.1, 53.6, 53.4, 48.6, 36.2, 30.1, 25.4, 24.3, 21.0, 14.1; MS (ESI+) for C₂₅H₂₇Cl₂N₃O₆S *m/z* 568.0 (M+H)⁺; MS (ESI-) for C₂₅H₂₇Cl₂N₃O₆S *m/z* 565.9 (M-H)⁻; MS (FAB) *m/z* (rel. intensity) 568 (MH⁺, 86), 644 (18), 571 (19), 570 (61), 569 (30), 568 (86), 335 (16), 188 (99), 173 (19), 141 (53), 116 (23); HRMS (FAB) calcd for C₂₅H₂₇Cl₂N₃O₆S+H₁ 568.1075, found 568.1096; Anal. Calcd

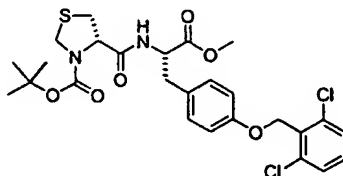
for $C_{25}H_{27}Cl_2N_3O_6S \cdot 0.4 H_2O$: C, 52.16; H, 4.87; N, 7.30. Found: C, 52.46; H, 4.90; N, 7.15. % Water (KF): 1.25.

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Example 32.

[S-(R*,R*)]-4-[[[1-[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is *t*-butyl, Y is CO_2 -, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*, *S*)).

10



Example 32 was prepared as described in Scheme A from D-cysteine using di-*t*-butyl dicarbonate to form the requisite carbamate. Physical data as follows: IR (liq.) 1745, 1702, 1565, 1511, 1467, 1439, 1368, 1299, 1241, 1197, 1177, 1162, 1017, 778, 768 cm^{-1} ; 1H NMR ($CDCl_3$) δ 7.36 (2 H), 7.25 (1 H), 7.06 (2 H), 6.94 (2 H), 5.23 (2 H), 4.75 (3 H), 4.12 (1 H), 3.72 (3 H), 3.37 (1 H), 3.14 (3 H), 1.45 (9 H); ^{13}C NMR ($CDCl_3$) δ 171.5, 157.9, 136.9, 132.0, 130.3, 128.4, 128.2, 115.0, 81.9, 65.1, 62.8, 53.2, 52.3, 49.2, 36.9, 28.1, 27.9; MS (ESI+) for $C_{26}H_{30}Cl_2N_2O_6S$ m/z 568.9 ($M+H$) $^+$; MS (ESI-) for $C_{26}H_{30}Cl_2N_2O_6S$ m/z 566.7 ($M-H$) $^-$; Anal. Calcd for $C_{26}H_{30}Cl_2N_2O_6S \cdot 0.09 H_2O$: C, 54.68; H, 5.33; N, 4.91. Found: C, 54.62; H, 5.41; N, 4.73. % Water (KF): 0.28.

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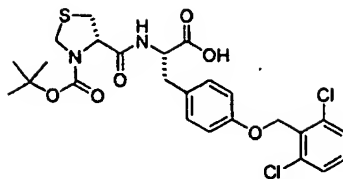
Example 33.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester

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(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is *t*-butyl, Y is CO_2 -, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*, *S*)).

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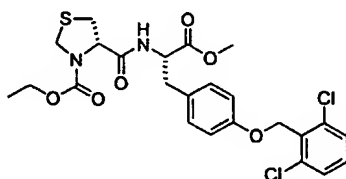


Example 33 was prepared from example 32 by the procedure described in preparation 6.

- Physical data as follows: IR (mull) 1734, 1704, 1676, 1612, 1565, 1511, 1439, 1393,
 1300, 1241, 1196, 1178, 1162, 777, 769 cm^{-1} ; ^1H NMR ($\text{DMSO}-d_6$) δ 8.24 (1 H), 7.50
 (3 H), 7.13 (2 H), 6.93 (2 H), 5.16 (2 H), 4.52 (3 H), 4.21 (1 H), 2.91 (4 H), 1.31 (9 H);
 ^{13}C NMR ($\text{DMSO}-d_6$) δ 172.7, 169.7, 157.1, 152.6, 135.9, 131.7, 131.5, 130.2, 129.8,
 128.7, 114.2, 79.8, 64.8, 61.6, 53.2, 49.2, 36.0, 34.8, 27.8, 21.0; MS (ESI+) for
 $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_2\text{O}_6\text{S}$ m/z 554.9 (M+H) $^+$; MS (ESI-) for $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_2\text{O}_6\text{S}$ m/z 552.8 (M-H) $^-$;
 Anal. Calcd for $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_2\text{O}_6\text{S} \cdot 0.15 \text{H}_2\text{O}$: C, 53.79; H, 5.11; N, 5.02. Found: C, 54.17;
 H, 5.17; N, 5.00. % Water (KF): 0.50.

Example 34.

- [S-(R*,R*)]-4-[[[1-[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-
 oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester
 (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is ethyl, Y is CO_2^- ,
 R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).



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Example was prepared as described in Scheme A from D-cysteine using ethyl
 chloroformate to form the requisite carbamate. Physical data as follows: IR (mull) 3282,
 1742, 1705, 1692, 1664, 1562, 1509, 1436, 1352, 1343, 1236, 1196, 1175, 1015, 786 cm^{-1} ;
 ^1H NMR (CDCl_3) δ 7.37 (2 H), 7.25 (1 H), 7.05 (2 H), 6.94 (2 H), 6.74 (1 H), 5.23 (2 H),

- 4.77 (3 H), 4.34 (1 H), 4.18 (2 H), 3.74 (3 H), 3.37 (1 H), 3.13 (3 H), 1.31 (3 H); ^{13}C NMR (CDCl_3) δ 171.6, 171.4, 158.0, 137.0, 132.1, 130.5, 130.4, 128.5, 128.2, 115.0, 65.2, 63.6, 63.2, 62.7, 53.6, 53.2, 52.4, 36.9, 14.5; MS (ESI+) for $\text{C}_{24}\text{H}_{26}\text{Cl}_2\text{N}_2\text{O}_6\text{S}$ m/z 540.9 ($\text{M}+\text{H}$) $^+$; HRMS (EI) calcd for $\text{C}_{24}\text{H}_{26}\text{Cl}_2\text{N}_2\text{O}_6\text{S}$ 540.0889, found 540.0878; Anal. Calcd for $\text{C}_{24}\text{H}_{26}\text{Cl}_2\text{N}_2\text{O}_6\text{S} \cdot 0.26 \text{ H}_2\text{O}$: C, 52.79; H, 4.89; N, 5.13. Found: C, 52.41; H, 4.82; N, 4.96. % Water (KF): 0.85.

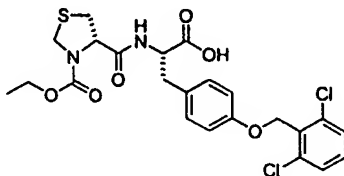
10

Example 35.

[S-(R*,R*)]-4-[[[1-carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester

(Scheme A, A-8: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is ethyl, Y is CO_2^- ,

- 15 R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).



Example 35 was prepared from example 34 by the procedure described in preparation 6.

- 20 Physical data as follows: IR (mull) 1709, 1675, 1612, 1565, 1511, 1439, 1416, 1346, 1300, 1241, 1196, 1179, 1115, 1018, 768 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.34 (2 H), 7.23 (1 H), 7.12 (2 H), 6.94 (2 H), 6.85 (1 H), 5.22 (2 H), 4.77 (4 H), 4.34 (1 H), 4.16 (2 H), 3.33 (4 H), 1.26 (3 H); ^{13}C NMR (CDCl_3) δ 174.0, 170.2, 158.1, 155.1, 137.0, 132.0, 130.5, 128.5, 128.1, 115.0, 65.2, 63.9, 63.0, 62.9, 53.3, 36.4, 21.9, 14.5; MS (ESI+) for $\text{C}_{23}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_6\text{S}$ m/z 527.0 ($\text{M}+\text{H}$) $^+$; MS (ESI-) for $\text{C}_{23}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_6\text{S}$ m/z 524.9 ($\text{M}-\text{H}$) $^-$; HRMS (EI) calcd for $\text{C}_{23}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_6\text{S}$ 526.0732, found 526.0726; Anal. Calcd for
- 25

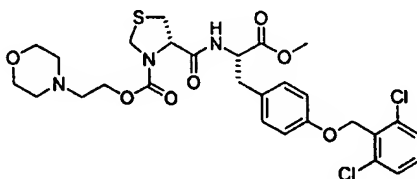
-63-

$C_{23}H_{24}Cl_2N_2O_6S \cdot 0.20 H_2O$: C, 52.02; H, 4.63; N, 5.27. Found: C, 52.12; H, 4.73; N, 5.34.
% Water (KF): 0.69.

Example 36.

[S-(R*,R*)]-4-[[[1-[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(4-morpholinyl)ethyl] ester

(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 2-(4-morpholinyl)ethyl, Y is CO_2^- , R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).



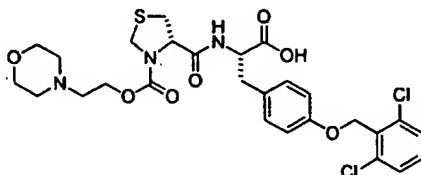
Example 36 was prepared as described in Scheme A from D-cysteine using 4-(2-hydroxyethyl)morpholine to form the requisite carbamate. Physical data as follows: mp 138-140°C; IR (mull) 3286, 1743, 1705, 1660, 1559, 1513, 1435, 1428, 1302, 1245, 1226, 1215, 1176, 1015, 764 cm^{-1} ; 1H NMR (300 MHz, DMSO- d_6) δ 8.43 (1 H), 7.48 (3 H), 7.13 (2 H), 6.94 (2 H), 5.16 (2 H), 4.59 (2 H), 4.48 (1 H), 4.26 (1 H), 4.07 (2 H), 3.63 (3 H), 3.51 (4 H), 3.23 (1 H), 3.01 (1 H), 2.84 (1 H), 2.71 (1 H), 2.41 (6 H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 172.2, 157.7, 153.9, 136.5, 132.2, 132.0, 130.8, 130.1, 129.2, 114.8, 66.6, 65.4, 63.4, 57.0, 53.9, 53.8, 52.4, 36.4; MS (ESI+) for $C_{28}H_{33}Cl_2N_3O_7S$ m/z 625.8 (M+H) $^+$; Anal. Calcd for $C_{28}H_{33}Cl_2N_3O_7S$: C, 53.67; H, 5.31; N, 6.71. Found: C, 53.69; H, 5.27; N, 6.69.

Example 37.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(4-morpholinyl)ethyl] ester

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(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 2-(4-morpholinyl)ethyl, Y is CO_2^- , R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).



5

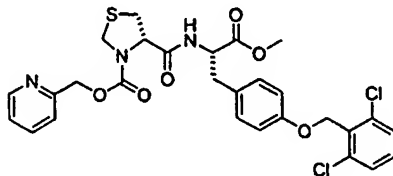
Example 37 was prepared from example 36 by the procedure described in preparation 6. Physical data as follows: IR (mull) 1710, 1610, 1585, 1565, 1511, 1439, 1408, 1351, 1301, 1240, 1196, 1179, 1116, 1017, 767 cm^{-1} ; ^1H NMR (300 MHz, DMSO-d_6) δ 8.27 (1 H), 7.55 (2 H), 7.45 (1 H), 7.13 (2 H), 6.94 (2 H), 5.16 (2 H), 4.59 (2 H), 4.40 (1 H), 4.27 (1 H), 4.04 (2 H), 3.52 (4 H), 3.21 (1 H), 2.86 (3 H), 2.44 (6 H), ^{13}C NMR (75 MHz, DMSO-d_6) δ 173.2, 172.5, 157.6, 136.5, 132.2, 132.0, 130.8, 130.4, 129.2, 114.7, 66.5, 65.3, 63.2, 57.0, 53.9, 53.7, 36.5, 21.5; MS (ESI+) for $\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 611.9 (M+H) $^+$; Anal. Calcd for $\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_7\text{S} \cdot 1.0 \text{ C}_2\text{H}_4\text{O}_2 \cdot 0.63 \text{ H}_2\text{O} \cdot 0.28 \text{ HCl}$: C, 50.13; H, 5.31; N, 6.03; Cl, 11.59. Found: C, 49.80; H, 5.30; N, 6.05; Cl, 11.20. % Water (KF): 1.58.

15

Example 38.

[S-(R*, R*)]-4-[[[1-[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[(2-pyridinyl)methyl] ester (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 2-pyridinylmethyl, Y is CO_2^- , R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).

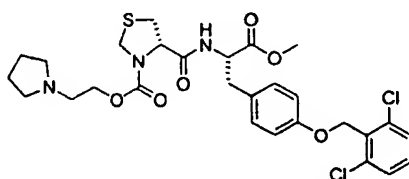
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Example 38 was prepared as described in Scheme A from D-cysteine using 2-pyridinemethanol to form the requisite carbamate. Physical data as follows: mp 123-125°C; IR (mull) 3334, 1728, 1709, 1668, 1531, 1511, 1441, 1405, 1345, 1294, 1286, 1236, 1228, 1015, 762 cm^{-1} ; ^1H NMR (300 MHz, DMSO- d_6) δ 8.57 (2 H), 7.79 (1 H), 7.54 (2 H), 7.42 (1 H), 7.27 (2 H), 7.12 (2 H), 6.92 (2 H), 5.13 (4 H), 4.69 (2 H), 4.49 (1 H), 4.34 (1 H), 3.59 (3 H), 3.24 (1 H), 2.89 (3 H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 172.2, 169.9, 157.7, 156.5, 153.6, 137.3, 136.5, 132.2, 132.0, 130.8, 130.1, 129.2, 123.2, 121.0, 114.8, 67.7, 65.3, 62.0, 54.0, 52.4, 50.3, 36.3, 35.3; HRMS (EI) calcd for $\text{C}_{28}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ 603.0997, found 603.0992; Anal. Calcd for $\text{C}_{28}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$: C, 55.63; H, 4.50; N, 6.95. Found: C, 55.56; H, 4.59; N, 6.93.

Example 39.

[S-(R*,R*)]-4-[[[1-[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(1-pyrrolidinyl)ethyl] ester (Scheme A, A-7: where $R_{A,1}$ and $R_{A,2}$ are the same and equal to H, R_3 is 2-(1-pyrrolidinyl)ethyl, Y is CO_2^- , R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).

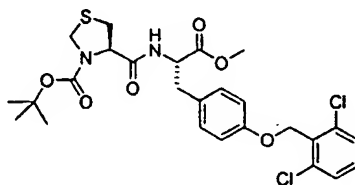


Example 39 was prepared as described in Scheme A from D-cysteine using 1-(2-hydroxyethyl)pyrrolidine to form the requisite carbamate. Physical data as follows: mp 130-132°C; IR (mull) 1745, 1702, 1661, 1556, 1513, 1435, 1426, 1303, 1245, 1226, 1214, 1176, 1017, 825, 765 cm^{-1} ; ^1H NMR (300 MHz, DMSO- d_6) δ 8.42 (1 H), 7.48 (3 H), 7.13 (2 H), 6.94 (2 H), 5.16 (2 H), 4.59 (2 H), 4.48 (1 H), 4.27 (1 H), 3.99 (2 H), 3.63 (3 H), 3.24 (1 H), 2.68 (9 H), 1.62 (4 H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 172.2, 170.0, 157.7, 136.5, 132.2, 132.0, 130.8, 130.1, 129.2, 114.8, 65.3, 65.1, 54.4, 53.9, 52.42, 36.35, 23.6;

MS (ESI+) for $C_{28}H_{33}Cl_2N_3O_6S$ m/z 609.8 (M+H)⁺; Anal. Calcd for $C_{28}H_{33}Cl_2N_3O_6S$: C, 55.08; H, 5.45; N, 6.88. Found: C, 54.72; H, 5.58; N, 6.60.

Example 40.

[R-(R*,S*)]-4-[[[1-[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester
(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is *t*-butyl, Y is CO₂-, R_4 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (R, S)).



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Example 40 was prepared as described in Scheme A from L-cysteine using di-*t*-butyl dicarbonate to form the requisite carbamate. Physical data as follows: IR (mull) 1746, 1702, 1611, 1565, 1511, 1439, 1299, 1241, 1197, 1177, 1162, 1118, 1016, 777, 768 cm⁻¹; ¹H NMR (CDCl₃) δ 7.39 (2 H), 7.25 (1 H), 6.93 (2 H), 6.93 (2 H), 5.24 (2 H), 4.72 (3 H), 4.20 (1 H), 3.74 (3 H), 3.35 (1 H), 3.12 (3 H), 1.44 (9 H); ¹³C NMR (CDCl₃) δ 171.7, 170.0, 169.5, 158.3, 137.3, 132.4, 130.7, 130.6, 128.8, 128.5, 115.2, 115.1, 82.4, 65.5, 53.5, 52.6, 50.3, 37.3, 28.4; MS (ESI+) for $C_{26}H_{30}Cl_2N_2O_6S$ m/z 554.9 (M+H)⁺; MS (ESI-) for $C_{26}H_{30}Cl_2N_2O_6S$ m/z 552.8 (M-H)⁻; Anal. Calcd for $C_{26}H_{30}Cl_2N_2O_6S \cdot 0.1 H_2O$: C, 54.65; H, 5.33; N, 4.90. Found: C, 54.59; H, 5.30; N, 4.88. % Water (KF): 0.33.

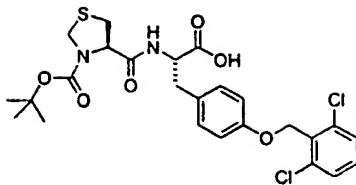
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Example 41.

[R-(R*,S*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester
(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is *t*-butyl, Y is CO₂-, R_4 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (R, S)).

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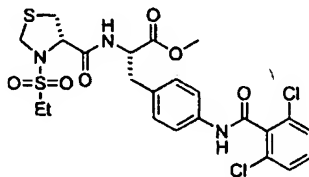


Example 41 was prepared from example 40 by the procedure described in preparation 6.

- 5 Physical data as follows: IR (mull) 1737, 1705, 1679, 1612, 1565, 1512, 1439, 1300, 1241, 1196, 1178, 1163, 1117, 777, 769 cm^{-1} ; ^1H NMR (DMSO-d_6) δ 8.07 (1 H), 7.54 (2 H), 7.44 (1 H), 7.16 (2 H), 6.93 (2 H), 5.16 (2 H), 4.56 (1 H), 4.39 (3 H), 3.56 (1 H), 2.84 (3 H), 1.23 (9 H);
- ^{13}C NMR (DMSO-d_6) δ 172.7, 169.9, 157.0, 152.6, 135.9, 131.6, 131.4, 130.1, 129.9, 129.8, 128.7, 114.1, 79.8, 66.9, 64.7, 61.7, 53.5, 49.3, 35.6, 34.8, 27.9, 27.6; MS (ESI+) for $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_2\text{O}_6\text{S}$ m/z 554.9 ($\text{M}+\text{H}^+$); MS (ESI-) for $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_2\text{O}_6\text{S}$ m/z 552.8 ($\text{M}-\text{H}^-$);
- 10 Anal. Calcd for $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_2\text{O}_6\text{S} \cdot 0.27 \text{H}_2\text{O}$: C, 53.59; H, 5.13; N, 5.00. Found: C, 53.97; H, 5.14; N, 4.96. % Water (KF): 0.86.

Example 42.

- 15 4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-(ethylsulfonyl)-4-thiazolidinyl]carbonyl]-L-phenylalanine methyl ester
- (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is ethyl, Y is SO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



20

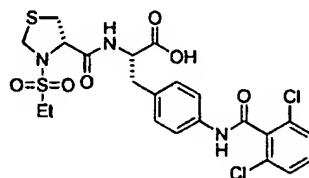
Example 42 was prepared as described in Scheme A from D-cysteine using ethanesulfonyl chloride to form the requisite sulfonamide. Physical data as follows: IR (mull) 1743, 1666, 1605, 1561, 1535, 1515, 1432, 1413, 1328, 1269, 1219, 1195, 1146, 799, 782 cm^{-1} ;

¹H NMR (CDCl₃) δ 7.58 (2 H), 7.46 (1 H), 7.32 (3 H), 7.17 (2 H), 7.07 (1 H), 4.85 (1 H), 4.73 (2 H), 4.28 (1 H), 3.76 (3 H), 3.54 (1 H), 3.26 (1 H), 3.05 (4 H), 1.40 (3 H); ¹³C NMR (CDCl₃) δ 171.3, 168.2, 162.4, 136.4, 135.8, 132.4, 132.3, 131.0, 130.1, 128.2, 120.6, 64.9, 53.2, 52.6, 51.5, 45.8, 37.1, 34.2, 31.0, 29.3, 7.7; MS (ESI+) for C₂₃H₂₅Cl₂N₃O₆S₂ *m/z* 573.9 (M+H)⁺; MS (ESI-) for C₂₃H₂₅Cl₂N₃O₆S₂ *m/z* 571.7 (M-H)⁻; HRMS (FAB) calcd for C₂₃H₂₅Cl₂N₃O₆S₂+H₁ 574.0640, found 574.0634; Anal. Calcd for C₂₃H₂₅Cl₂N₃O₆S₂ • 0.1 H₂O: C, 47.97; H, 4.40; N, 7.30. Found: C, 48.36; H, 4.59; N, 6.80.

Example 43.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-(ethylsulfonyl)-4-thiazolidinyl]carbonyl]-L-phenylalanine

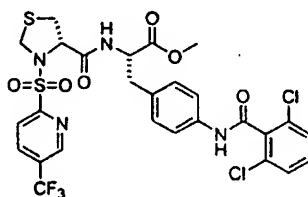
(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is ethyl, Y is SO₂, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



Example 43 was prepared from example 42 by the procedure described in preparation 6.. Physical data as follows: IR (mull) 1734, 1664, 1605, 1562, 1536, 1516, 1432, 1414, 1330, 1272, 1234, 1195, 1146, 799, 781 cm⁻¹; ¹H NMR (DMSO-d₆) δ 12.91 (1 H), 10.65 (1 H), 8.15 (1 H), 7.51 (5 H), 7.17 (2 H), 4.77 (2 H), 4.43 (1 H), 4.29 (1 H), 3.94 (6 H), 1.20 (3 H); ¹³C NMR (DMSO-d₆) δ 172.5, 168.9, 161.9, 137.1, 136.5, 133.2, 131.4, 131.3, 129.8, 128.3, 119.3, 63.7, 53.6, 51.4, 45.3, 36.1, 34.7; MS (ESI+) for C₂₂H₂₃Cl₂N₃O₆S₂ *m/z* 559.9 (M+H)⁺; MS (ESI-) for C₂₂H₂₃Cl₂N₃O₆S₂ *m/z* 557.8 (M-H)⁻; HRMS (FAB) calcd for C₂₂H₂₃Cl₂N₃O₆S₂+H₁ 560.0483, found 560.0488; Anal. Calcd for C₂₂H₂₃Cl₂N₃O₆S₂ • 0.72 H₂O: C, 46.08; H, 4.30; N, 7.33. Found: C, 46.42; H, 4.37; N, 7.01. % Water (KF): 2.26.

Example 44.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-[[5-(trifluoromethyl)-2-pyridinyl]sulfonyl]-4-thiazolidinyl]carbonyl]-L-phenylalanine methyl ester
 (Scheme A, A-7: where $R_{A,1}$ and $R_{A,2}$ are the same and equal to H, R_3 is 2-(5-trifluoromethylpyridyl), Y is SO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).

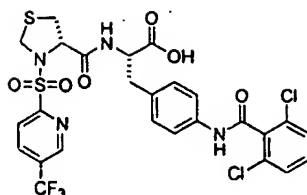


Example 44 was prepared as described in Scheme A from D-cysteine using 2-(5-trifluoromethylpyridyl)sulfonyl chloride to form the requisite sulfonamide. Physical data as follows: IR (mull) 1745, 1668, 1603, 1535, 1515, 1432, 1413, 1327, 1219, 1179, 1142, 1108, 1073, 1016, 616 cm^{-1} ; 1H NMR ($CDCl_3$) δ 8.76 (1 H), 8.17 (2 H), 7.90 (1 H), 7.51 (2 H), 7.32 (4 H), 7.17 (2 H), 5.18 (1 H), 4.96 (1 H), 4.66 (1 H), 4.31 (1 H), 3.78 (3 H), 3.52 (1 H), 3.15 (3 H); ^{13}C NMR ($CDCl_3$) δ 171.3, 168.4, 162.2, 147.2, 136.2, 136.0, 132.4, 131.0, 130.3, 128.2, 123.0, 120.4, 120.3, 65.7, 53.6, 52.5, 51.4, 37.3, 34.0; MS (ESI+) for $C_{27}H_{23}Cl_2F_3N_4O_6S_2$ m/z 690.8 (M+H) $^+$; MS (ESI+) for $C_{27}H_{23}Cl_2F_3N_4O_6S_2$ m/z 712.9 (M+Na) $^+$; Anal. Calcd for $C_{27}H_{23}Cl_2F_3N_4O_6S_2 \cdot 0.2 H_2O$: C, 46.68; H, 3.39; N, 8.06. Found: C, 46.60; H, 3.52; N, 7.92. % Water (KF): 0.47.

Example 45.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-[[5-(trifluoromethyl)-2-pyridinyl]sulfonyl]-4-thiazolidinyl]carbonyl]-L-phenylalanine
 (Scheme A, A-8: where $R_{A,1}$ and $R_{A,2}$ are the same and equal to H, R_3 is 2-(5-trifluoromethylpyridyl), Y is SO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).

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Example 45 was prepared from example 44 by the procedure described in preparation 6.

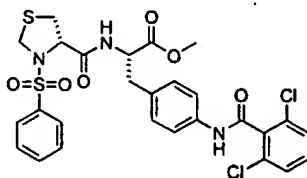
Physical data as follows: IR (mull) 1740, 1666, 1602, 1562, 1533, 1517, 1432, 1354,

- 5 1327, 1179, 1143, 1108, 1074, 1016, 613 cm^{-1} ; ^1H NMR (DMSO-d_6) δ 10.63 (1 H), 9.24 (1 H), 8.54 (1 H), 8.46 (1 H), 8.18 (1 H), 7.50 (5 H), 7.17 (2 H), 5.00 (1 H), 4.74 (1 H), 4.42 (2 H), 3.04 (2 H), 2.90 (1 H), 2.78 (1 H); ^{13}C NMR (DMSO-d_6) δ 172.8, 168.9, 162.3, 159.0, 147.8, 137.5, 137.4, 136.9, 133.5, 131.8, 131.7, 130.2, 128.7, 123.8, 119.7, 64.7, 53.9, 52.3, 36.7, 35.1; MS (ESI+) for $\text{C}_{26}\text{H}_{21}\text{Cl}_2\text{F}_3\text{N}_4\text{O}_6\text{S}_2$ m/z 676.5 (M+H) $^+$; MS (ESI-) for
- 10 $\text{C}_{26}\text{H}_{21}\text{Cl}_2\text{F}_3\text{N}_4\text{O}_6\text{S}_2$ m/z 674.5 (M-H) $^-$; Anal. Calcd for $\text{C}_{26}\text{H}_{21}\text{Cl}_2\text{F}_3\text{N}_4\text{O}_6\text{S}_2 \cdot 0.33$: C, 45.69; H, 3.20; N, 8.20. Found: C, 45.81; H, 3.38; N, 8.13. % Water (KF): 0.88.

Example 46.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-(phenylsulfonyl)-4-thiazolidinyl]carbonyl]-L-phenylalanine methyl ester

- 15 (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is phenyl, Y is SO_2 , R_4 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



- 20 Example 46 was prepared as described in Scheme A from D-cysteine using benzenesulfonyl chloride to form the requisite sulfonamide. Physical data as follows: IR (mull) 1744, 1668, 1604, 1531, 1515, 1432, 1413, 1355, 1324, 1268, 1220, 1195, 1167, 1090, 730 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.84 (2 H), 7.65 (5 H), 7.45 (1 H), 7.30 (6 H), 4.90 (1 H), 4.63 (2 H), 4.37 (1 H), 3.75 (3 H), 3.32 (1 H), 3.15 (2 H), 2.53 (1 H); ^{13}C NMR

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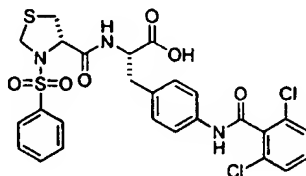
(CDCl₃) δ 171.2, 168.2, 162.4, 136.4, 136.3, 134.1, 132.4, 131.0, 130.2, 129.6, 128.2, 128.1, 127.9, 120.6, 65.3, 53.3, 52.6, 51.8, 37.4, 33.3; MS (ESI+) for C₂₇H₂₃Cl₂N₃O₆S₂ m/z 621.8 (M+H)⁺; MS (ESI-) for C₂₇H₂₃Cl₂N₃O₆S₂ m/z 619.8 (M-H)⁻; Anal. Calcd for C₂₇H₂₃Cl₂N₃O₆S₂ • 0.2 H₂O: C, 51.84; H, 4.18; N, 6.72. Found: C, 51.72; H, 4.18; N, 6.52.

5 % Water (KF): 0.48.

Example 47.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-(phenylsulfonyl)-4-thiazolidinyl]carbonyl]-L-phenylalanine

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is phenyl, Y is SO₂,
10 R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



Example 47 was prepared from example 46 by the procedure described in preparation 6.

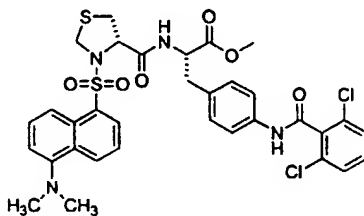
15 Physical data as follows: IR (mull) 1735, 1666, 1605, 1562, 1533, 1516, 1432, 1414, 1352, 1328, 1195, 1180, 1167, 1090, 731 cm⁻¹; ¹H NMR (CDCl₃) δ 8.11 (1 H), 7.82 (2 H), 7.66 (3 H), 7.56 (2 H), 7.23 (6 H), 4.91 (1 H), 4.66 (1 H), 4.60 (1 H), 4.35 (1 H), 3.30 (1 H), 3.19 (2 H), 2.59 (1 H); ¹³C NMR (DMSO-d₆) δ 172.2, 168.2, 161.7, 137.0, 136.9, 136.3, 133.6, 133.0, 131.2, 131.1, 129.6, 129.3, 128.1, 127.6, 119.2, 63.9, 53.4, 51.5, 48.4,
20 36.0, 33.8; MS (ESI+) for C₂₆H₂₃Cl₂N₃O₆S₂ m/z 607.9 (M+H)⁺; MS (FAB) m/z (rel. intensity) 608 (MH⁺, 85), 610 (67), 608 (85), 466 (30), 371 (41), 228 (38), 193 (38), 149 (30), 129 (31), 118 (99), 63 (35); HRMS (FAB) calcd for C₂₆H₂₃Cl₂N₃O₆S₂+H₁ 608.0483, found 608.0491; Anal. Calcd for C₂₆H₂₃Cl₂N₃O₆S₂ • 0.27 H₂O: C, 50.91; H, 3.87; N, 6.85. Found: C, 50.68; H, 4.05; N, 6.65. % Water (KF): 0.79.

25

Example 48.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-[[5-(dimethylamino)-1-naphthalenyl]sulfonyl]-4-thiazolidinyl]carbonyl]-L-phenylalanine methyl ester

(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 5-dimethylamino-1-naphthyl, Y is SO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



5

Example 48 was prepared as described in Scheme A from D-cysteine using 5-dimethylamino-1-naphthalenesulfonyl chloride to form the requisite sulfonamide. Physical data as follows: IR (mull) 1744, 1684, 1605, 1562, 1533, 1515, 1431, 1412, 1350, 1324, 1231, 1202, 1163, 1145, 798 cm^{-1} ; 1H NMR ($CDCl_3$) δ 8.64 (1 H), 8.34 (2 H), 7.53 (5 H), 7.29 (4 H), 6.87 (3 H), 4.93 (1 H), 4.75 (1 H), 4.64 (1 H), 4.31 (1 H), 3.69 (3 H), 3.47 (1 H), 2.84 (8 H), 2.46 (1 H); ^{13}C NMR ($CDCl_3$) δ 171.2, 167.7, 162.5, 136.3, 135.9, 132.4, 132.3, 131.9, 131.5, 131.0, 130.1, 129.7, 129.2, 128.2, 124.4, 120.4, 65.1, 53.3, 52.5, 50.1, 45.9, 37.1, 33.3; MS (ESI+) for $C_{33}H_{32}Cl_2N_4O_6S_2$ m/z 736.8 ($M+Na$) $^+$; Anal. Calcd for $C_{33}H_{32}Cl_2N_4O_6S_2 \cdot 0.17 H_2O$: C, 55.15; H, 4.54; N, 7.79. Found: C, 55.20; H, 4.73; N, 7.49. % Water (KF): 0.43.

15

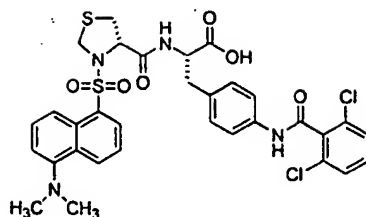
Example 49.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-[[5-(dimethylamino)-1-naphthalenyl]sulfonyl]-4-thiazolidinyl]carbonyl]-L-phenylalanine

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 5-dimethylamino-1-naphthyl, Y is SO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).

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Example 49 was prepared from example 48 by the procedure described in preparation 6.

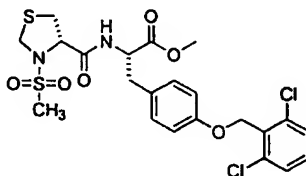
Physical data as follows: IR (mull) 1666, 1605, 1587, 1577, 1562, 1532, 1516, 1431,

- 5 1412, 1395, 1325, 1163, 1145, 798, 631 cm^{-1} ; ^1H NMR (DMSO-d_6) δ 10.60 (1 H), 8.54 (1 H), 8.28 (2 H), 7.54 (8 H), 7.25 (1 H), 6.93 (2 H), 4.95 (1 H), 4.84 (1 H), 4.39 (1 H), 4.17 (1 H), 2.95 (2 H), 2.80 (7 H), 2.54 (1 H); ^{13}C NMR (DMSO-d_6) δ 167.2, 161.7, 151.4, 136.6, 136.3, 133.6, 133.1, 131.1, 130.8, 130.3, 129.5, 129.3, 129.0, 128.8, 128.1, 123.6, 118.9, 118.3, 115.3, 63.6, 54.2, 50.3, 44.9, 36.7, 33.9, 21.0; MS (ESI+) for
- 10 $\text{C}_{32}\text{H}_{30}\text{Cl}_2\text{N}_4\text{O}_6\text{S}_2$ m/z 700.8 ($\text{M}+\text{H}^+$); HRMS (FAB) calcd for $\text{C}_{32}\text{H}_{30}\text{Cl}_2\text{N}_4\text{O}_6\text{S}_2+\text{H}_1$ 701.1062, found 701.1039

Example 50.

O-[(2,6-Dichlorophenyl)methyl]-N-[[[(4S)-3-(methylsulfonyl)-4-thiazolidinyl]carbonyl]-L-tyrosine methyl ester

- 15 (Scheme A, A-7: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is methyl, Y is SO_2 , R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).



- 20 Example 50 was prepared as described in Scheme A from D-cysteine using methanesulfonyl chloride to form the requisite sulfonamide. Physical data as follows: IR (mull) 1742, 1680, 1611, 1564, 1510, 1439, 1345, 1299, 1240, 1179, 1158, 1016, 976, 779, 768 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.36 (2 H), 7.24 (1 H), 7.08 (3 H), 6.97 (2 H), 5.25 (2 H), 4.77 (3 H), 4.29 (1 H), 3.74 (3 H), 3.43 (1 H), 3.53 (1 H), 3.10 (2 H), 2.93 (3 H);

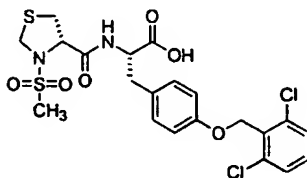
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^{13}C NMR (CDCl_3) δ 171.4, 168.0, 158.2, 137.0, 132.1, 130.3, 128.5, 127.9, 115.3, 65.2, 65.0, 59.4, 53.5, 53.3, 52.5, 51.9, 42.2, 37.3, 36.9, 34.2; MS (ESI+) for $\text{C}_{22}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_6\text{S}_2$ m/z 546.8 ($\text{M}+\text{H}$) $^+$; MS (ESI+) for $\text{C}_{22}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_6\text{S}_2$ m/z 568.8 ($\text{M}+\text{H}$) $^+$; HRMS (EI) calcd for $\text{C}_{22}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_6\text{S}_2$ 546.0453, found 546.0448; Anal. Calcd for $\text{C}_{22}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_6\text{S}_2 \cdot 0.07$ H_2O : C, 48.15; H, 4.43; N, 5.10. Found: C, 48.17; H, 4.51; N, 5.02. % Water (KF): 0.24.

Example 51.

O-[(2,6-Dichlorophenyl)methyl]-N-[(4S)-3-(methylsulfonyl)-4-thiazolidinyl]carbonyl]-L-tyrosine

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is methyl, Y is SO_2 , R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).



Example 51 was prepared from example 50 by the procedure described in preparation 6.

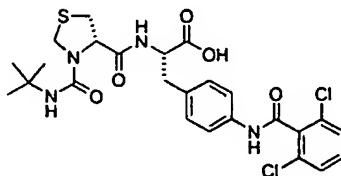
Physical data as follows: IR (mull) 1737, 1675, 1611, 1565, 1511, 1439, 1345, 1300, 1241, 1197, 1179, 1157, 1016, 778, 769 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.34 (2 H), 7.16 (5 H), 6.99 (2 H), 5.23 (2 H), 4.85 (1 H), 4.68 (2 H), 4.27 (1 H), 3.51 (1 H), 3.32 (1 H), 3.15 (2 H), 2.93 (3 H); ^{13}C NMR (CDCl_3) δ 174.8, 168.7, 158.3, 137.0, 132.1, 130.5, 130.4, 128.5, 127.6, 115.4, 115.2, 65.2, 64.9, 53.1, 52.0, 37.1, 36.4, 34.3; MS (ESI+) for $\text{C}_{21}\text{H}_{22}\text{Cl}_2\text{N}_2\text{O}_6\text{S}_2$ m/z 532.8 ($\text{M}+\text{H}$) $^+$; MS (ESI-) for $\text{C}_{21}\text{H}_{22}\text{Cl}_2\text{N}_2\text{O}_6\text{S}_2$ m/z 530.7 ($\text{M}-\text{H}$) $^-$; HRMS (FAB) calcd for $\text{C}_{21}\text{H}_{22}\text{Cl}_2\text{N}_2\text{O}_6\text{S}_2+\text{H}$ 533.0374, found 533.0386; Anal. Calcd for $\text{C}_{21}\text{H}_{22}\text{Cl}_2\text{N}_2\text{O}_6\text{S}_2 \cdot 0.06 \text{H}_2\text{O}$: C, 47.19; H, 4.17; N, 5.24. Found: C, 47.58; H, 4.35; N, 5.10. % Water (KF): 0.20.

Example 52.

4-[(2,6-Dichlorobenzoyl)amino]-N-[(4S)-3-[(1,1-dimethylethyl)amino]carbonyl]-4-thiazolidinyl]carbonyl]-L-phenylalanine

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(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is *t*-butyl, Y is CONH-, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*, *S*)).



5

Example 52 was prepared from example 6 by the procedure described in preparation 6.

Physical properties as follows: IR (mull) 3289, 1728, 1664, 1607, 1580, 1561, 1536,

1432, 1414, 1394, 1326, 1270, 1242, 1213, 1195 cm^{-1} ; ^1H NMR (300 MHz, CD_3OD) δ

7.47 (2 H), 7.24 (3 H), 7.04 (2 H), 4.63 (2 H), 4.37 (1 H), 4.17 (1 H), 3.09 (4 H), 1.22

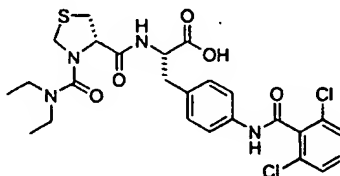
10 (9 H); ^{13}C NMR (75 MHz, CDCl_3) δ 176.9, 174.8, 167.2, 159.9, 140.5, 140.0, 136.4, 136.1, 134.6, 133.8, 131.8, 124.5, 66.6, 57.1, 55.3, 53.0, 40.1, 37.1, 32.9; MS (ESI+) for $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_4\text{O}_5\text{S}$ m/z 566.9 ($\text{M}+\text{H}^+$), 588.9 ($\text{M}+\text{Na}^+$); MS (ESI+) for $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_4\text{O}_5\text{S}$ m/z 566.9 ($\text{M}+\text{H}^+$); HRMS (FAB) calcd for $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_4\text{O}_5\text{S} + \text{H}^+$ 567.1235, found 567.1253.

Example 53.

15 4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4*S*)-3-[(diethylamino)carbonyl]-4-thiazolidinyl]carbonyl]-L-phenylalanine

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is ethyl, Y is $\text{CON}(\text{CH}_2\text{CH}_3)-$, R_5 is 4-[(2,6-dichlorobenzoyl)amino] phenyl and stereochemistry is (*S*, *S*)).

20



Example 53 was prepared from example 7 by the procedure described in preparation 6.

Physical properties as follows: IR (mull) 3269, 1734, 1663, 1607, 1562, 1535, 1515,

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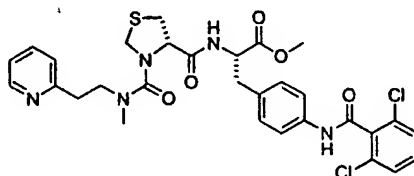
1431, 1415, 1348, 1325, 1269, 1213, 1195, 799 cm^{-1} ; ^1H NMR (300 MHz, CD_3OD) δ 7.57 (2 H), 7.35 (3 H), 7.09 (2 H), 5.09 (1 H), 4.76 (1 H), 4.38 (2 H), 3.31 (3 H), 3.13 (5 H), 1.05 (6 H); ^{13}C NMR (75 MHz, CD_3OD) δ 176.6, 173.9, 167.1, 166.3, 140.7, 140.1, 136.1, 134.6, 133.8, 131.8, 124.1, 68.5, 57.4, 56.6, 45.9, 40.4, 36.6, 16.8; MS (ESI+) for

5 $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_4\text{O}_5\text{S}$ m/z 567.1 ($\text{M}+\text{H}^+$); Anal. Calcd for $\text{C}_{25}\text{H}_{28}\text{Cl}_2\text{N}_4\text{O}_5\text{S}$: C, 52.91; H, 4.97; N, 9.87.

Found: C, 52.60; H, 5.13; N, 9.47.

Example 54.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-[[methyl[2-(2-
 10 pyridinyl)ethyl]amino]carbonyl]-4-thiazolidinyl]carbonyl]-L-phenylalanine methyl ester
 (Scheme A, A-7: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is 2-(2-pyridyl)ethyl,
 Y is $\text{CON}(\text{CH}_3)-$, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S,
 S)).



15

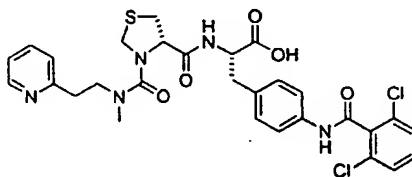
Example 54 was prepared as described in Scheme A from D-cysteine using 2-(2-methylaminoethyl)pyridine to form the requisite urea. Physical data as follows: mp 80-90°C (dec); IR (mull) 1743, 1665, 1606, 1561, 1538, 1514, 1489, 1432, 1413, 1395, 1323,
 1268, 1216, 1195, 799 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 8.44 (1 H), 8.34 (1 H),
 20 7.66 (1 H), 7.51 (5 H), 7.19 (4 H), 4.72 (1 H), 4.48 (1 H), 4.40 (1 H), 4.20 (1 H), 3.58
 (4 H), 3.42 (1 H), 3.89 (10 H); ^{13}C NMR (75 MHz, $\text{DMSO}-d_6$) δ 172.1, 170.3, 162.3,
 161.8, 159.4, 149.4, 137.6, 137.0, 136.8, 133.3, 131.8, 131.7, 130.0, 128.7, 123.8, 122.0,
 119.8, 64.58, 53.8, 52.7, 52.5, 49.7, 36.5, 36.2, 35.7, 33.4; MS (ESI+) for $\text{C}_{30}\text{H}_{31}\text{Cl}_2\text{N}_5\text{O}_5\text{S}$
 25 m/z 643.9 ($\text{M}+\text{H}^+$); Anal. Calcd for $\text{C}_{30}\text{H}_{31}\text{Cl}_2\text{N}_5\text{O}_5\text{S}$: C, 55.90; H, 4.85; N, 10.86. Found:
 C, 55.52; H, 5.09; N, 10.64.

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Example 55.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-[[methyl[2-(2-pyridinyl)ethyl]amino]carbonyl]-4-thiazolidinyl]carbonyl]-L-phenylalanine

5 (Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 2-(2-pyridyl)ethyl, Y is $\text{CON}(\text{CH}_3)-$, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



10

Example 55 was prepared from example 54 by the procedure described in preparation 6.

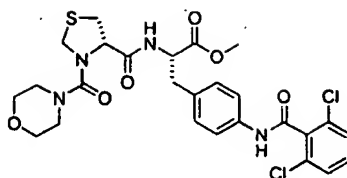
Physical data as follows: IR (mull) 1682, 1656, 1606, 1561, 1540, 1513, 1432, 1413, 1398, 1323, 1268, 1242, 1195, 799, 780 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 10.66 (1 H), 8.45 (1 H), 8.09 (1 H), 7.66 (1 H), 7.51 (5 H), 7.20 (4 H), 4.72 (1 H), 4.40 (2 H),
 15 4.23 (1 H), 3.61 (1 H), 3.40 (1 H), 3.04 (7 H), 2.79 (3 H); MS (ESI+) for $\text{C}_{29}\text{H}_{29}\text{Cl}_2\text{N}_5\text{O}_5\text{S}$ m/z 629.9 ($\text{M}+\text{H}^+$); Anal. Calcd for $\text{C}_{29}\text{H}_{29}\text{Cl}_2\text{N}_5\text{O}_5\text{S} \cdot 0.61 \text{ H}_2\text{O}$: C, 54.29; H, 4.75; N, 10.92. Found: C, 54.29; H, 5.00; N, 10.32. % Water (KF): 1.72.

Example 56.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4S)-3-(4-morpholinylcarbonyl)-4-thiazolidinyl]carbonyl]-L-phenylalanine methyl ester

20

(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 and Y together form CO-morpholino, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).

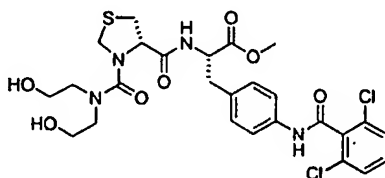


Example 56 was prepared as described in Scheme A from D-cysteine using morpholine to form the requisite urea. Physical data as follows: mp 223-225°C; ¹H NMR (300 MHz, DMSO-d₆) δ 10.66 (1 H), 8.25 (1 H), 7.51 (5 H), 7.16 (2 H), 4.81 (1 H), 4.60 (1 H), 4.50 (1 H), 4.28 (1 H), 3.64 (3 H), 3.53 (4 H), 3.09 (8 H); ¹³C NMR (75 MHz, DMSO-d₆) 172.1, 170.3, 162.3, 161.4, 137.6, 136.8, 133.3, 131.8, 131.7, 130.0, 128.7, 119.8, 66.2, 64.3, 53.8, 52.8, 52.5, 46.8, 36.1, 33.7; HRMS (FAB) calcd for C₂₆H₂₈Cl₂N₄O₆S+H₁ 595.1185, found 595.1189; Anal. Calcd for C₂₆H₂₈Cl₂N₄O₆S: C, 52.44; H, 4.74; N, 9.41. Found: C, 52.42; H, 4.96; N, 9.23.

Example 57.

N-[[[(4S)-3-[[Bis(2-hydroxyethyl)amino]carbonyl]-4-thiazolidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine methyl ester

15 (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is 2-hydroxyethyl and Y is $\text{CON}(\text{CH}_2\text{CH}_2\text{OH})$, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



20

Example 57 was prepared as described in Scheme A from D-cysteine using diethanolamine to form the requisite urea. Physical data as follows: mp 105-107°C; IR (mull) 3284, 1743, 1662, 1608, 1561, 1539, 1516, 1432, 1414, 1355, 1326, 1270, 1217, 1196, 799 cm^{-1} ; ^1H NMR (300 MHz, DMSO- d_6) δ 10.69 (1 H), 8.28 (1 H), 7.56 (4 H),

7.47 (1 H), 7.16 (2 H), 4.87 (3 H), 4.66 (1 H), 4.47 (1 H), 4.25 (1 H), 3.63 (3 H), 3.47 (6 H), 3.02 (6 H); MS (ESI+) for $C_{26}H_{30}Cl_2N_4O_7S$ m/z 612.9 (M+H)⁺; Anal. Calcd for $C_{26}H_{30}Cl_2N_4O_7S \cdot 0.47 H_2O$: C, 50.21; H, 5.01; N, 9.01. Found: C, 50.02; H, 5.00; N, 8.93. % Water (KF): 1.36.

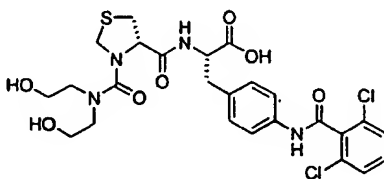
5

Example 58.

N-[[[(4S)-3-[[Bis(2-hydroxyethyl)amino]carbonyl]-4-thiazolidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_1 is 2-hydroxyethyl and Y is $CON(CH_2CH_2OH)$, R_2 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and

10 stereochemistry is (S, S)).



Example 58 was prepared from example 57 by the procedure described in preparation 6.

15 Physical data as follows: IR (mull) 3281, 3196, 1724, 1660, 1608, 1580, 1561, 1542, 1515, 1431, 1415, 1354, 1328, 1271, 1196 cm^{-1} ; 1H NMR (300 MHz, DMSO- d_6) δ 10.65 (1 H), 8.13 (1 H), 7.55 (4 H), 7.47 (1 H), 7.16 (2 H), 4.85 (2 H), 4.66 (1 H), 4.40 (1 H), 4.27 (1 H), 3.48 (6 H), 3.01 (6 H); MS (ESI+) for $C_{25}H_{28}Cl_2N_4O_7S$ m/z 598.9 (M+H)⁺; Anal. Calcd for $C_{25}H_{28}Cl_2N_4O_7S \cdot 1.04 H_2O$: C, 58.58; H, 4.90; N, 9.06. Found: C, 48.88; 20 H, 5.05; N, 8.79. % Water (KF): 3.02.

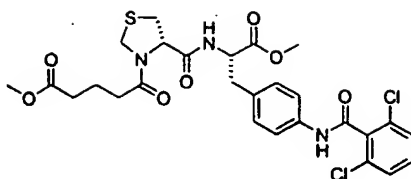
Example 59.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]- δ -oxo-3-thiazolidinepentanoic acid 3-methyl ester

(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to H, R_1 is

25 $CH_2CH_2CH_2CO_2CH_3$, Y is CO, R_2 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)).

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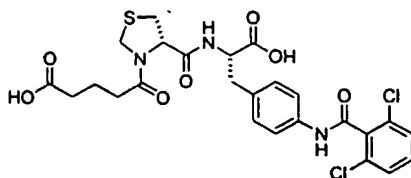


Example 59 was prepared as described in Scheme A from D-cysteine using methyl glutaryl chloride to form the requisite amide. Physical data as follows: IR (mull) 3266, 1741, 1734, 1685, 1678, 1630, 1610, 1560, 1545, 1441, 1435, 1414, 1327, 1268, 1227 cm^{-1} ; ^1H NMR ($\text{DMSO}-d_6$) δ 10.65 (1 H), 8.43 (1 H), 7.50 (5 H), 7.15 (2 H), 4.72 (2 H), 4.44 (2 H), 3.63 (3 H), 3.56 (3 H), 3.10 (4 H), 2.15 (4 H), 1.70 (2 H); ^{13}C NMR (CDCl_3) δ 171.6, 170.6, 170.1, 169.9, 169.6, 161.9, 137.2, 136.5, 132.9, 131.3, 129.7, 128.3, 119.4, 61.4, 53.7, 53.5, 52.0, 51.3, 48.7, 36.4, 35.8, 35.1, 33.1, 32.8, 32.5, 25.4, 19.8; MS (ESI+) for $\text{C}_{27}\text{H}_{29}\text{N}_3\text{O}_7\text{S}$ m/z 610.0 ($\text{M}+\text{H}^+$); MS (ESI-) for $\text{C}_{27}\text{H}_{29}\text{N}_3\text{O}_7\text{S}$ m/z 608.0 ($\text{M}-\text{H}^-$); Anal. Calcd for $\text{C}_{27}\text{H}_{29}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$: C, 53.12; H, 4.79; N, 6.88. Found: C, 52.81; H, 4.90; N, 6.88

Example 60.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]- δ -oxo-3-thiazolidinepentanoic acid

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is $\text{CH}_2\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$, Y is CO, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)).



Example 60 was prepared from example 59 by the procedure described in preparation 12. Physical data as follows: IR (mull) 3271, 3193, 3124, 1725, 1661, 1607, 1561, 1539,

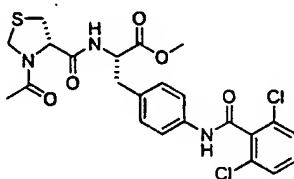
-81-

1516, 1432, 1414, 1327, 1271, 1195, 799 cm^{-1} ; ^1H NMR ($\text{DMSO}-d_6$) δ 12.40 (1 H), 10.63 (1 H), 8.31 (1 H), 7.50 (5 H), 7.16 (2 H), 4.75 (2 H), 4.34 (2 H), 2.95 (4 H), 2.15 (4 H), 1.66 (2 H); ^{13}C NMR (CD_3CN) δ 175.0, 172.7, 172.6, 171.0, 163.5, 137.8, 137.0, 134.4, 132.7, 132.3, 131.1, 130.9, 129.1, 120.7, 63.2, 54.5, 49.9, 37.2, 34.1, 33.3, 20.7; MS (ESI+) for $\text{C}_{25}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 582.0 ($\text{M}+\text{H}$) $^+$; MS (ESI-) for $\text{C}_{25}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 579.9 ($\text{M}-\text{H}$) $^-$; Anal. Calcd for $\text{C}_{25}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_7\text{S} \cdot \text{H}_2\text{O}$: C, 50.01; H, 4.53; N, 7.00. Found: C, 49.61; H, 4.38; N, 6.61.

Example 61.

10 N-[[[(4S)-3-Acetyl-4-thiazolidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine methyl ester

(Scheme A, A-7: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is CH_3 , Y is CO, R_4 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)).



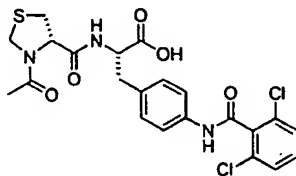
15

Example 61 was prepared as described in Scheme A from D-cysteine using acetyl chloride to form the requisite amide. Physical data as follows: IR (mull) 3260, 3067, 1748, 1686, 1623, 1608, 1561, 1542, 1515, 1445, 1429, 1419, 1324, 1267, 1221 cm^{-1} ; ^1H NMR ($\text{DMSO}-d_6$) δ 8.44 (1 H), 7.52 (5 H), 7.17 (2 H), 4.78 (2 H), 4.37 (2 H), 3.63 (3 H), 3.06 (4 H), 1.94 (3 H); ^{13}C NMR ($\text{DMF}-d_7$) δ 172.4, 170.6, 138.4, 137.5, 133.8, 132.3, 131.9, 130.6, 128.9, 120.0, 70.9, 63.6, 62.4, 54.7, 52.3, 50.2, 49.8, 49.2, 37.2, 36.6, 22.7; MS (ESI+) for $\text{C}_{23}\text{H}_{23}\text{Cl}_2\text{N}_3\text{O}_5\text{S}$ m/z 523.9 ($\text{M}+\text{H}$) $^+$; MS (ESI+) for $\text{C}_{23}\text{H}_{23}\text{Cl}_2\text{N}_3\text{O}_5\text{S}$ m/z 545.8 ($\text{M}+\text{Na}$) $^+$; HRMS (FAB) calcd for $\text{C}_{23}\text{H}_{23}\text{Cl}_2\text{N}_3\text{O}_5\text{S}+\text{H}$, 524.0814, found 524.0812; Anal. Calcd for $\text{C}_{23}\text{H}_{23}\text{Cl}_2\text{N}_3\text{O}_5\text{S} \cdot 0.1 \text{H}_2\text{O}$: C, 52.46; H, 4.45; N, 7.98. Found: C, 52.85; H, 4.42; N, 8.00. % Water (KF): 0.24.

Example 62.

N-[[[(4S)-3-Acetyl-4-thiazolidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_1 is CH_3 , Y is CO, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)).



Example 62 was prepared from example 61 by the procedure described in preparation 6.

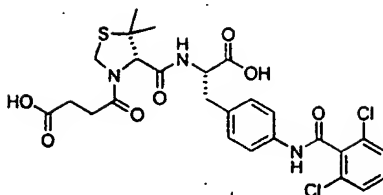
Physical data as follows: IR (mull) 3070, 1747, 1682, 1663, 1625, 1608, 1580, 1561, 1548, 1514, 1443, 1431, 1416, 1278, 1220 cm^{-1} ; ^1H NMR ($\text{DMSO}-d_6$) δ 12.48 (1 H), 10.63 (1 H), 8.29 (1 H), 7.50 (5 H), 4.73 (2 H), 4.34 (2 H), 2.97 (4 H), 1.93 (3 H); ^{13}C NMR ($\text{DMSO}-d_6$) δ 172.4, 169.6, 169.2, 168.5, 168.0, 161.7, 136.9, 136.3, 133.2, 133.0, 131.2, 129.6, 128.1, 119.2, 62.2, 61.0, 53.5, 53.3, 49.3, 48.4, 36.3, 35.7, 35.0, 33.1, 22.4, 20.9; MS (ESI-) for $\text{C}_{22}\text{H}_{21}\text{Cl}_2\text{N}_3\text{O}_5\text{S}$ m/z 507.9 (M-H)⁺; Anal. Calcd for $\text{C}_{22}\text{H}_{21}\text{Cl}_2\text{N}_3\text{O}_5\text{S} \cdot 0.1 \text{ H}_2\text{O}$: C, 51.57; H, 4.18; N, 8.20. Found: C, 51.49; H, 4.36; N, 8.07. % Water (KF): 0.40.

Example 63.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-5,5-dimethyl- δ -oxo-3-thiazolidinebutanoic acid

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to CH_3 , R_3 is $\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$, Y is CO, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (S,S)).

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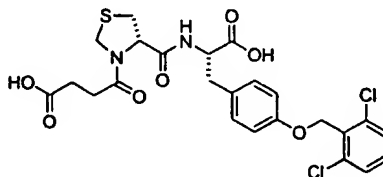


Example 63 was prepared as described in Scheme A from D-penicillamine using methyl succinyl chloride to form the requisite amide. Physical data as follows: IR (mull) 3264, 3198, 3071, 1721, 1660, 1608, 1562, 1541, 1516, 1432, 1415, 1327, 1270, 1195, 799 cm^{-1} ; ^1H NMR (DMSO- d_6) δ 10.62 (1 H), 8.15 (1 H), 7.52 (5 H), 7.18 (2 H), 4.72 (2 H), 4.43 (3 H), 2.91 (4 H), 2.17 (2 H), 1.36 (3 H), 1.07 (3 H); ^{13}C NMR (DMSO- d_6) δ 173.5, 172.6, 170.2, 169.7, 168.4, 168.1, 161.7, 136.9, 136.3, 133.3, 131.1, 129.6, 129.4, 128.1, 119.2, 70.6, 53.8, 53.5, 51.6, 48.4, 47.7, 35.9, 30.7, 30.5, 28.9, 28.6, 24.1; MS (ESI+) for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 596.0 (M+H) $^+$; MS (ESI+) for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 617.9 (M+Na) $^+$. MS (ESI-) for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 593.8 (M-H) $^-$; MS (FAB) m/z (rel. intensity) 596 (MH $^+$, 20), 598 (15), 596 (20), 331 (11), 193 (13), 141 (15), 139 (99), 116 (16), 107 (13), 105 (50), 89 (25); HRMS (FAB) calcd for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S}+\text{H}$, 596.1025, found 596.1036.

Example 64.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]carbonyl]- γ -oxo-3-thiazolidinebutanoic acid

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is $\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$, Y is CO, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, and stereochemistry is (S,S)).



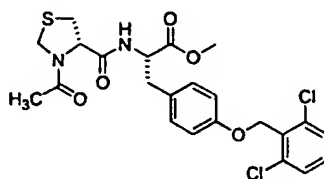
Example 64 was prepared as described in Scheme A from D-cysteine using methyl succinyl chloride to form the requisite amide. Physical data as follows: IR (mull) 3073,

3031, 1725, 1640, 1612, 1585, 1565, 1535, 1511, 1439, 1300, 1241, 1196, 1179, 768 cm^{-1} ;
 ^1H NMR ($\text{DMSO}-d_6$) δ 8.26 (1 H), 7.53 (2 H), 7.44 (1 H), 7.12 (2 H), 6.93 (2 H), 5.16
 (2 H), 4.75 (2 H), 4.40 (2 H), 4.20 (1 H), 2.81 (4 H), 2.32 (2 H), 2.07 (1 H); ^{13}C NMR
 ($\text{DMSO}-d_6$) δ 173.8, 172.6, 170.0, 169.3, 157.1, 136.0, 131.5, 130.4, 130.0, 129.8, 128.7,
 114.2, 64.8, 61.5, 53.6, 48.5, 36.1, 35.5, 35.0, 33.1, 28.9, 28.6, 21.0; MS (ESI+) for
 $\text{C}_{24}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_7\text{S}$ m/z 554.8 ($\text{M}+\text{H}$) $^+$; MS (ESI-) for $\text{C}_{24}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_7\text{S}$ m/z 552.7 ($\text{M}-\text{H}$) $^-$;
 HRMS (FAB) calcd for $\text{C}_{24}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_7\text{S}+\text{H}$, 555.0759, found 555.0750.

Example 65.

N-[[[(4S)-3-Acetyl-4-thiazolidinyl]carbonyl]-O-[(2,6-dichlorophenyl)methyl]-L-
 tyrosine methyl ester

(Scheme A, A-7: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is CH_3 , Y is CO, R_4
 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, and stereochemistry is (S,S)).



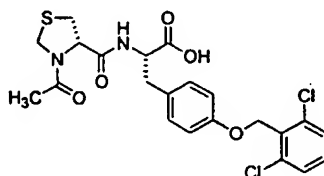
15

Example 65 was prepared as described in Scheme A from D-cysteine using acetyl chloride
 to form the requisite amide. Physical data as follows: IR (mull) 1744, 1657, 1612, 1585,
 1564, 1511, 1438, 1405, 1352, 1299, 1240, 1197, 1179, 1016, 768 cm^{-1} ; ^1H NMR (CDCl_3)
 δ 7.36 (2 H), 7.24 (1 H), 6.97 (5 H), 5.24 (2 H), 5.04 (1 H), 4.78 (1 H), 4.50 (2 H), 3.74
 (3 H), 3.45 (1 H), 3.17 (3 H), 2.02 (3 H); ^{13}C NMR (CDCl_3) δ 172.7, 171.6, 171.4, 169.8,
 168.9, 168.7, 157.8, 136.9, 131.9, 130.3, 129.9, 128.6, 128.4, 114.9, 65.1, 61.6, 56.0,
 53.6, 53.3, 49.6, 36.7, 31.7, 22.5; MS (ESI+) for $\text{C}_{23}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_7\text{S}$ m/z 532.9 ($\text{M}+\text{Na}$) $^+$; MS
 (EI) m/z (rel. intensity) 510 (M^+ , 1), 338 (42), 337 (12), 336 (63), 267 (12), 265 (18), 163
 (10), 161 (63), 159 (99), 130 (9), 88 (43); Anal. Calcd for $\text{C}_{23}\text{H}_{24}\text{Cl}_2\text{N}_2\text{O}_7\text{S}\cdot 0.19 \text{H}_2\text{O}$: C,
 53.66; H, 4.77; N, 5.44. Found: C, 53.81; H, 4.75; N, 5.33. % Water (KF): 0.66.

Example 66.

N-[[[(4S)-3-Acetyl-4-thiazolidinyl]carbonyl]-O-[(2,6-dichlorophenyl)methyl]-L-tyrosine

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is CH_3 , Y is CO, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, and stereochemistry is (S,S)).



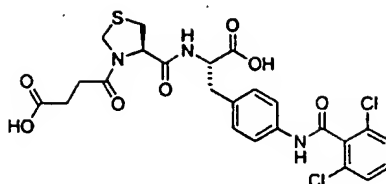
Example 66 was prepared from example 65 by the procedure described in preparation 6.

Physical data as follows: IR (mull) 1730, 1646, 1612, 1585, 1565, 1511, 1439, 1414, 1299, 1240, 1196, 1179, 1016, 779, 768 cm^{-1} ; 1H NMR (DMSO- d_6) δ 8.26 (1 H), 7.54 (2 H), 7.44 (1 H), 7.13 (2 H), 6.92 (2 H), 5.16 (2 H), 4.72 (2 H), 4.31 (2 H), 2.91 (5 H), 1.92 (3 H); ^{13}C NMR (DMSO- d_6) δ 172.6, 169.2, 168.0, 157.1, 136.0, 131.7, 131.5, 130.3, 128.7, 114.2, 64.8, 62.2, 61.0, 53.7, 49.4, 48.5, 36.1, 35.5, 35.1, 33.2, 22.4, 21.0; MS (ESI+) for $C_{22}H_{22}Cl_2N_2O_5S$ m/z 496.9 (M+H) $^+$; MS (ESI-) for $C_{22}H_{22}Cl_2N_2O_5S$ m/z 494.8 (M-H) $^-$; MS (FAB) m/z (rel. intensity) 497 (MH $^+$, 99), 617 (29), 573 (12), 539 (25), 500 (11), 499 (78), 498 (38), 497 (99), 496 (11), 225 (62), 130 (14); HRMS (FAB) calcd for $C_{22}H_{22}Cl_2N_2O_5S+H$, 497.0705, found 497.0713; Anal. Calcd for $C_{22}H_{22}Cl_2N_2O_5S \cdot 0.41 H_2O$: C, 52.35; H, 4.56; N, 5.55. Found: C, 52.65; H, 4.51; N, 5.50. % Water (KF): 1.46.

Example 67.

[R-(R*,S*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]- γ -oxo-3-thiazolidinebutanoic acid (Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is $CH_2CH_2CO_2H$, Y is CO, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (R,S)).

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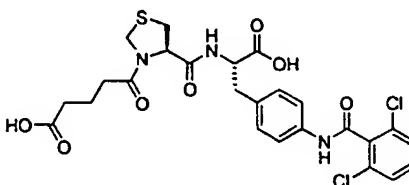
Example 67 was prepared as described in Scheme A from L-cysteine using methyl succinyl chloride to form the requisite amide. Physical data as follows: ¹H NMR

(CD₃CN) δ 8.86 (1 H), 7.55 (2 H), 7.42 (3 H), 7.24 (2 H), 7.11 (1 H), 4.90 (1 H), 4.65 (2 H), 4.33 (1 H), 3.14 (3 H), 2.47 (6 H), 1.80 (1 H); ¹³C NMR (CD₃CN) δ 174.7, 172.6, 172.3, 170.7, 163.4, 137.7, 137.0, 134.4, 132.7, 132.2, 130.9, 129.1, 63.2, 60.9, 54.4, 49.7, 37.1, 36.7, 32.7, 30.6, 30.0, 29.5, 21.1, 14.4; MS (FAB) *m/z* (rel. intensity) 568 (MH⁺, 99), 646 (11), 644 (16), 572 (13), 571 (12), 570 (73), 569 (38), 568 (99), 567 (15), 216 (22), 88 (27); Anal. Calcd for C₂₄H₂₃Cl₂N₃O₇S • 0.5 H₂O: C, 49.92; H, 4.19; N, 7.23. Found: C, 50.01; H, 4.54; N, 7.05.

Example 68.

[R-(R*,S*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-δ-oxo-3-thiazolidinepentanoic acid

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R₃ is CH₂CH₂CH₂CO₂H, Y is CO, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (R,S)).



Example 67 was prepared as described in Scheme A from L-cysteine using methyl glutaryl chloride to form the requisite amide. Physical data as follows: ¹H NMR (CD₃CN) δ 8.85 (1 H), 7.55 (2 H), 7.44 (3 H), 7.17 (3 H), 4.86 (1 H), 4.64 (2 H), 4.34 (1 H), 3.64 (1 H),

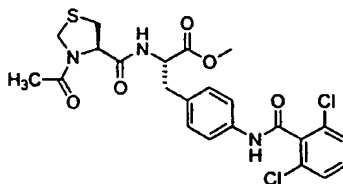
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3.13 (3 H), 2.27 (4 H), 1.79 (4 H); ^{13}C NMR (CD_3CN) δ 175.0, 172.9, 172.7, 170.6, 163.4, 137.7, 137.0, 134.2, 132.7, 132.2, 131.0, 129.1, 120.7, 68.2, 63.1, 54.2, 49.8, 37.1, 36.9, 34.0, 33.2, 32.6, 26.2, 20.6; MS (FAB) m/z (rel. intensity) 582 (MH^+ , 99), 585 (12), 584 (68), 583 (35), 582 (99), 581 (11), 88 (23), 69 (8), 57 (9), 55 (11), 43 (11); Anal. Calcd for $\text{C}_{25}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_5\text{S} \cdot 0.2 \text{ H}_2\text{O}$: C, 51.24; H, 4.37; N, 7.17. Found: C, 51.25; H, 4.68; N, 6.92.

Example 69.

N-[[[(4R)-3-Acetyl-4-thiazolidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine methyl ester

(Scheme A, A-7: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_1 is CH_3 , Y is CO, R_2 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (R,S)).



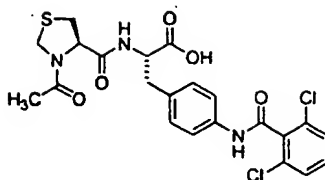
Example 69 was prepared as described in Scheme A from L-cysteine using acetyl chloride to form the requisite amide. Physical data as follows: IR (mull) 3268, 1743, 1662, 1607, 1561, 1538, 1515, 1431, 1413, 1354, 1324, 1270, 1217, 1195, 799 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.56 (3 H), 7.31 (4 H), 7.14 (2 H), 4.92 (2 H), 4.49 (1 H), 4.29 (1 H), 3.77 (3 H), 3.54 (1 H), 3.26 (1 H), 3.00 (2 H), 2.11 (3 H); ^{13}C NMR (CDCl_3) δ 171.5, 170.3, 168.9, 152.5, 135.9, 135.7, 132.9, 132.3, 131.0, 130.1, 129.9, 128.2, 128.0, 127.9, 120.4, 63.9, 61.4, 53.1, 52.5, 49.7, 37.0, 31.2, 22.6; MS (ESI+) for $\text{C}_{25}\text{H}_{23}\text{Cl}_2\text{N}_3\text{O}_5\text{S}$ m/z 523.8 ($\text{M}+\text{H}$) $^+$.

Example 70.

N-[[[(4R)-3-Acetyl-4-thiazolidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine

(Scheme A, A-8: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_1 is CH_3 , Y is CO, R_2 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (R,S)).

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Example 70 was prepared from example 69 by the procedure described in preparation 6.

Physical data as follows: IR (mull) 3276, 3068, 1727, 1661, 1607, 1561, 1540, 1516,

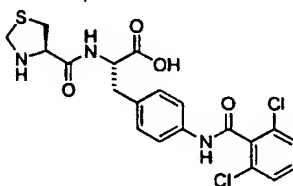
5 1444, 1431, 1414, 1327, 1271, 1218, 1195 cm^{-1} ; ^1H NMR (DMSO-d_6) δ 12.78 (1 H), 10.67 (1 H), 8.27 (1 H), 7.49 (4 H), 7.19 (2 H), 4.75 (2 H), 4.45 (3 H), 3.01 (3 H), 1.96 (3 H); ^{13}C NMR (DMSO-d_6) δ 172.6, 172.3, 169.4, 168.6, 168.2, 161.7, 136.9, 136.3, 133.3, 132.8, 131.1, 129.6, 128.1, 119.1, 62.2, 60.9, 53.4, 49.3, 48.7, 35.9, 35.1, 32.7, 22.4; MS (ESI+) for $\text{C}_{22}\text{H}_{21}\text{Cl}_2\text{N}_3\text{O}_5\text{S}$ m/z 509.8 ($\text{M}+\text{H}$) $^+$; MS (ESI-) for $\text{C}_{22}\text{H}_{21}\text{Cl}_2\text{N}_3\text{O}_5\text{S}$ m/z 507.8 (M-H) $^-$; HRMS (FAB) calcd for $\text{C}_{22}\text{H}_{21}\text{Cl}_2\text{N}_3\text{O}_5\text{S}+\text{H}$, 510.0657, found 510.0667;

Example 71.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4R)-4-thiazolidinyl]carbonyl]-L-phenylalanine monohydrochloride salt

(Scheme A, A-10: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to H, R_3 is 4-[(2,6-

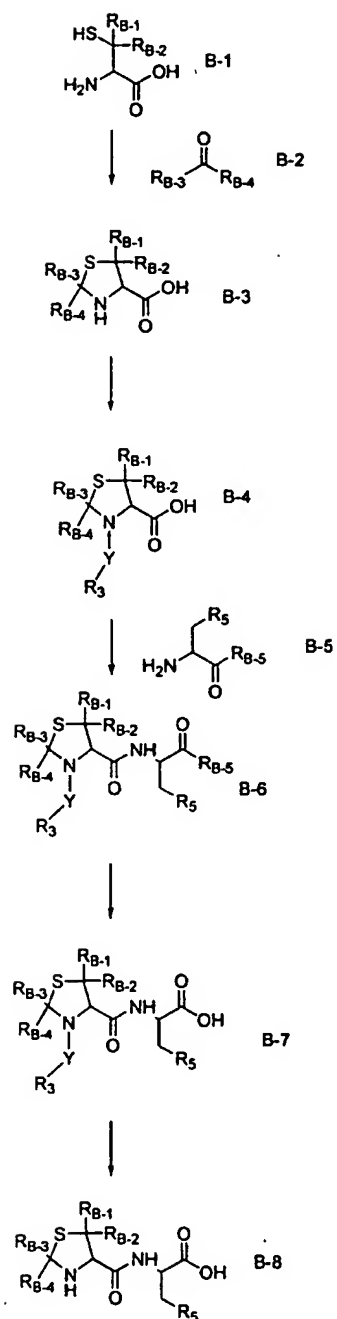
15 dichlorobenzoyl)amino]phenyl, and stereochemistry is (R,S)).



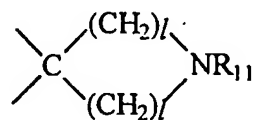
Example 71 was prepared as described in Scheme A from L-cysteine. Physical data as follows: IR (mull) 3249, 3190, 3036, 1729, 1662, 1605, 1578, 1562, 1541, 1516, 1432, 1414, 1328, 1271, 1195 cm^{-1} ; ^1H NMR (DMSO-d_6) δ 10.71 (1 H), 8.83 (1 H), 7.56 (5 H), 7.24 (2 H), 4.50 (1 H), 4.25 (1 H), 4.21 (2 H), 3.62 (1 H), 3.01 (4 H); ^{13}C NMR (DMSO-d_6) δ 172.0, 166.8, 161.8, 137.7, 132.7, 131.3, 131.0, 129.5, 128.1, 119.4, 72.0, 70.4, 62.3, 53.8, 49.0, 35.7, 33.4; MS (ESI+) for $\text{C}_{20}\text{H}_{19}\text{Cl}_2\text{N}_3\text{O}_4\text{S}$ m/z 468.1 ($\text{M}+\text{H}$) $^+$;

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Scheme B



where R_{B-1} , R_{B-2} , R_{B-3} , and R_{B-4} are defined as R_1 and in addition R_{B-3} and R_{B-4} may be attached to the same carbon atom and form a cyclic ring of 5-8 atoms of the formula:



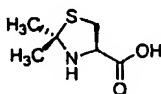
together with the carbon atom to which they are attached; R_{B-5} is defined as OH or O-(C_{1-6} alkyl)

Scheme B describes a general method for the preparation of thiazolidine-4-carboxylic acid derivatives of general structure **B-6**, **B-7** and **B-8** which are disubstituted at the 2 position (i.e., R_{B-3} and R_{B-4} are not equal to H). Within this class of structures, the nitrogen is derivatized immediately after forming the heterocyclic ring. Accordingly, a commercially available or readily prepared sulfur-containing amino acid of structure **B-1** (which is the same structure as **A-1**) is condensed with a commercially available or readily prepared ketone to afford the thiazolidine-4-carboxylic acid of general formula **B-3** (preparation 15) (for a general discussion of the condensation of aldehydes and ketones with cysteine or similar sulfur containing amino acids see: Coppola, G.M.; Schuster, H.F. *Asymmetric Synthesis: Construction of Chiral Molecules Using Amino Acids*; John Wiley: New York, 1987; Chapter 6, 171.). The amine group may be reacted with a variety of electrophilic reagents such as sulfonyl chlorides, carbonates, chloroformates, isocyanates, phosgene (or a suitable equivalent) and an amine, acid chlorides, and carboxylic acid anhydrides as described in Scheme A for the reaction of **A-6**. Preparation 16 is provided as a specific example of the synthesis of a compound of general structure **B-4**. Condensation of **B-4** with amino acid derivative **B-5** under standard peptide synthesis conditions provides the compound of general structure **B-6** (preparation 17). Mild base hydrolysis of the methyl ester of general structure **B-6** (where R_{B-5} is OCH_3) may be effected as described and exemplified in Scheme A (preparation 6 or 13) to afford compounds of the general structure **B-7**. Alternatively, in those cases where R_{B-5} is *O-t*-Bu, mild acidolysis can also provide compounds of the general structure **B-7** (by the procedure described in preparation 4 of Scheme A). In the case of *t*-butoxycarbonyl derivatised analogs of general structure **B-7** (i.e. where R_3 is *t*-butyl and Y is CO_2), mild

acidolysis affords compounds of general structure B-8 (by the procedure described in preparation 4 of Scheme A).

Preparation 15.

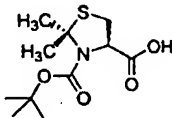
(Scheme B, B-3: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} are the same and equal to CH_3 , and stereochemistry is (*R*)).



A suspension of L-cysteine hydrochloride monohydrate (Scheme B, B-1: where R_{B-1} and R_{B-2} are the same and equal to H and stereochemistry is (*R*)) (20 g, 0.11 mol) in acetone (Scheme B, B-2: where R_{B-3} and R_{B-4} are equal to CH_3) (800 mL) was heated to reflux for 8 h. Cooling to room temperature resulted in precipitation of a solid which was collected by filtration, washed with acetone and dried *in vacuo* to afford the title compound (17.46 g) as a white solid: mp 165-167°C; (Lit. 165-168°C: Sheehan, J.C.; Yang, D-D.H. *J. Am. Chem. Soc.* 1957, 80, 1158) 1H NMR (D_2O) δ 4.75 (1 H), 3.59 (1 H), 3.44 (1 H), 1.73 (3 H), 1.71 (3 H); MS (ESI-) for $C_6H_{11}NO_2S$ m/z 159.9 (M-H).

Preparation 16.

(Scheme B, B-4: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} are the same and equal to CH_3 , Y is CO_2 , R_3 is *t*-butyl and stereochemistry is (*R*)).



To a solution of B-3 (Scheme B where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} are the same and equal to CH_3 , and stereochemistry is (*R*)) (17.46 g, 0.11 mol) in acetonitrile (250 mL) at ambient temperature was added di-*t*-butyl dicarbonate (25.64 g, 0.117 mol) followed by N,N-diisopropylethylamine (16.9 mL, 0.097 mol). The reaction mixture was stirred for 2 days and volatiles removed *in vacuo*. The residue was slurried in

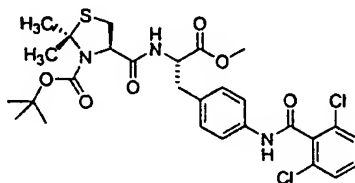
diethyl ether and filtered through a pad of celite. The filtrate was washed with 0.1 N HCl, water, brine, dried (Na_2SO_4), filtered and concentrated in vacuo. Crystallization of the clear oil from hexane provided the title compound (3.85 g) as a white solid: mp 125-126°C; (Lit. 114°C: Woodward, R.B.; Heusler, K.; Gosteli, J.; Naegeli, P.; Oppolzer, W.; Ramage, R.; Ranganathan, S.; Vorbruggen, H. *J. Am. Chem. Soc.* **1966**, *88*, 852) ^1H NMR (CDCl_3) δ 8.70 (1 H), 4.89 (1 H), 3.27 (2 H), 1.81 (6 H); MS (ESI-) for $\text{C}_{11}\text{H}_{19}\text{NO}_4\text{S}$ m/z 260.1 (M-H).

Preparation 17 and Example 72.

10 [R-(R*,S*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxo-ethyl]amino]carbonyl]-2,2-dimethyl-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl)ester

(Scheme B, B-6: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} are the same and equal to CH_3 , Y is CO_2 , R_3 is *t*-butyl, R_{B-5} is OCH_3 , R_5 is 4-[(2,6-

15 dichlorobenzoyl)amino]phenyl and stereochemistry is (*R*, *S*)).

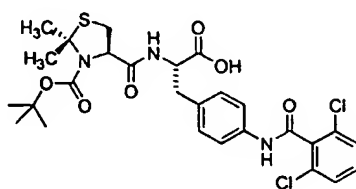


To a cooled (0-5°C) suspension of B-4 (Scheme B where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} are the same and equal to CH_3 , Y is CO_2 , R_3 is *t*-butyl and stereochemistry is (*R*)) (1.0 g, 3.83 mmol) and HOBT (638 mg, 4.17 mmol) in CH_2Cl_2 (20 mL) was added a solution of EDC (799 mg, 4.17 mmol) in CH_2Cl_2 (20 mL). After 30 min at 0-5°C, B-5 (Scheme B where R_{B-5} is OCH_3 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*)) (1.93 g, 4.77 mmol) was added followed by 4-methylmorpholine (520 μL , 4.77 mmol). The reaction mixture was gradually warmed to room temperature, stirred an additional 18 h and diluted with CH_2Cl_2 . The organic layer was separated and washed with 0.1 N HCl, sat. aqueous NaHCO_3 , brine, dried (Na_2SO_4),

filtered and concentrated in vacuo. Flash chromatography of the residue using CH₂Cl₂/acetone (3%) as eluant afforded the title compound (2.05 g) as a white solid: IR (mull) 1746, 1666, 1606, 1562, 1537, 1515, 1432, 1413, 1347, 1325, 1259, 1214, 1195, 1169, 799 cm⁻¹; ¹H NMR (CDCl₃) δ 7.54 (2 H), 7.32 (4 H), 7.20 (2 H), 6.90 (1 H), 4.78 (2 H), 3.74 (3 H), 3.17 (4 H), 1.74 (3 H), 1.43 (9 H); ¹³C NMR (CDCl₃) δ 171.6, 136.7, 136.2, 135.9, 132.7, 132.4, 132.3, 131.0, 130.6, 130.4, 130.0, 129.9, 128.2, 127.9, 120.5, 120.2, 120.1, 67.3, 53.4, 52.4, 42.8, 37.3, 36.9, 34.9, 30.9, 28.3; MS (ESI+) for C₂₈H₃₃Cl₂N₃O₆S *m/z* 610 (M+H)⁺; MS (ESI-) for C₂₈H₃₃Cl₂N₃O₆S *m/z* 607.9 (M-H)⁻; Anal. Calcd for C₂₈H₃₃Cl₂N₃O₆S • 0.13 H₂O: C, 54.88; H, 5.47; N, 6.86. Found: C, 54.66; H, 5.57; N, 6.73. % Water (KF): 0.37.

Example 73.

[R-(R*,S*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-2,2-dimethyl-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl)ester
(Scheme B, B-7: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} are the same and equal to CH₃, Y is CO₂, R₃ is *t*-butyl, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*R*, *S*)).



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Example 73 was prepared from example 72 by the procedure described in preparation 6. Physical data as follows: IR (mull) 1738, 1665, 1606, 1562, 1535, 1516, 1432, 1413, 1347, 1259, 1213, 1194, 1167, 799, 777 cm⁻¹; ¹H NMR (CD₃CN) δ 8.89 (1 H), 7.54 (2 H), 7.41 (3 H), 7.25 (2 H), 4.66 (2 H), 3.15 (4 H), 1.72 (3 H), 1.70 (3 H), 1.35 (9 H); ¹³C NMR (CD₃CN) δ 177.3, 171.8, 163.4, 137.9, 137.1, 133.9, 132.7, 132.3, 131.3, 129.1, 120.7, 81.2, 79.3, 68.0, 54.5, 37.4, 28.4; MS (ESI-) for C₂₇H₃₁Cl₂N₃O₆S *m/z* 593.9 (M-H)⁻; Anal.

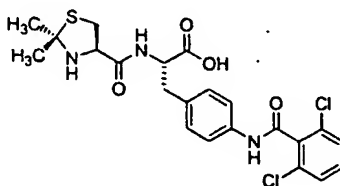
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Calcd for $C_{27}H_{31}Cl_2N_3O_6S \cdot 0.5 H_2O$: C, 53.56; H, 5.33; N, 6.94. Found: C, 53.77; H, 5.39; N, 6.70.

Example 74.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(4R)-2,2-dimethyl-4-thiazolidinyl]carbonyl]-L-phenylalanine

(Scheme B, B-8: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} are the same and equal to CH_3 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (R, S)).



Example 74 was prepared from example 73 by the procedure described in preparation 4.

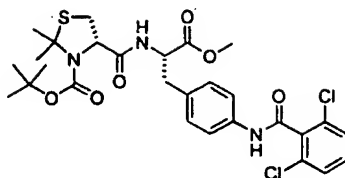
Physical data as follows: IR (mull) 3244, 3192, 3049, 1726, 1664, 1605, 1578, 1562, 1541, 1516, 1432, 1414, 1327, 1195, 799 cm^{-1} ; 1H NMR (DMSO- d_6) δ 13.00 (1 H), 10.69

(1 H), 7.52 (6 H), 7.24 (2 H), 4.52 (2 H), 3.12 (2 H), 2.93 (1 H), 1.62 (9 H); ^{13}C NMR (CD $_3$ OD) δ 166.9, 136.8, 136.0, 133.5, 131.8, 130.9, 129.5, 129.4, 127.9, 120.4, 120.2, 120.1, 60.8, 54.3, 54.0, 36.2, 36.0, 25.1; MS (ESI+) for $C_{22}H_{23}Cl_2N_3O_4S$ m/z 496.2 (M+H) $^+$; MS (ESI-) for $C_{22}H_{23}Cl_2N_3O_4S$ m/z 494.2 (M-H) $^-$; Anal. Calcd for $C_{22}H_{23}Cl_2N_3O_4S \cdot HCl \cdot 0.50 H_2O$: C, 48.76; H, 4.65; N, 7.75. Found: C, 48.56; H, 4.72; N, 7.49.

Example 75.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxo-ethyl]amino]carbonyl]-2,2-dimethyl-3-thiazolidinecarboxylic acid 3-(1,1-dimethylethyl)ester

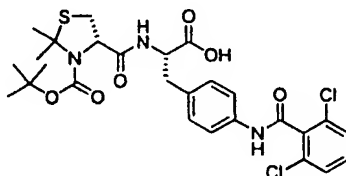
(Scheme B, B-6: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} are the same and equal to CH_3 , Y is CO_2 , R_3 is *t*-butyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



Example 75 was prepared as described in Scheme B from D-cysteine and acetone using di-
t-butyl dicarbonate to form the requisite carbamate. Physical data as follows: IR (mull)
 5 1745, 1686, 1666, 1605, 1537, 1515, 1432, 1413, 1349, 1325, 1259, 1213, 1206, 1195,
 1169 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.56 (2 H), 7.32 (3 H), 7.16 (2 H), 6.92 (1 H), 4.83 (2 H),
 3.73 (3 H), 3.20 (4 H), 1.78 (6 H), 1.45 (9 H); ^{13}C NMR (CDCl_3) δ 171.6, 170.0, 136.2,
 132.6, 132.4, 131.0, 130.9, 130.1, 129.9, 128.2, 120.5, 120.4, 67.3, 53.3, 52.5, 52.4, 37.4,
 28.4; MS (ESI+) for $\text{C}_{28}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 610.0 ($\text{M}+\text{H}^+$); MS (ESI-) for $\text{C}_{28}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$
 10 m/z 607.9 ($\text{M}-\text{H}^-$); Anal. Calcd for $\text{C}_{28}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$: C, 55.08; H, 5.45; N, 6.88; Cl, 11.61;
 S, 5.25. Found: C, 54.87; H, 5.47; N, 6.78

Example 76.

15 [S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-
 dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-2,2-dimethyl-3-
 thiazolidinecarboxylic acid 3-(1,1-dimethylethyl) ester
 (Scheme B, B-7: where $\text{R}_{\text{B-1}}$ and $\text{R}_{\text{B-2}}$ are the same and equal to H, $\text{R}_{\text{B-3}}$ and $\text{R}_{\text{B-4}}$ are the
 same and equal to CH_3 , Y is CO_2 , R_3 is *t*-butyl, R_5 is 4-[(2,6-
 20 dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*, *S*)).



Example 76 was prepared from example 75 by the procedure described in preparation 6.

Physical data as follows: IR (mull) 3280, 1739, 1665, 1606, 1562, 1535, 1516, 1432,

1413, 1348, 1272, 1259, 1195, 1167, 799 cm^{-1} ; ^1H NMR (CD_3CN) δ 8.83 (1 H), 7.55

(2 H), 7.43 (3 H), 7.22 (2 H), 6.83 (1 H), 4.68 (2 H), 3.07 (5 H), 1.73 (6 H), 1.40 (9 H);

5 ^{13}C NMR (CD_3CN) δ 172.6, 163.3, 137.9, 133.4, 133.3, 132.7, 132.1, 131.1, 130.2, 130.1, 129.1, 128.1, 128.0, 120.6, 119.5, 54.1, 37.4, 29.4, 28.5, 27.6; MS (ESI+) for

$\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 595.9 (M+H) $^+$; MS (ESI-) for $\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 593.7 (M-H) $^-$; MS

(FAB) m/z (rel. intensity) 596 (MH $^+$, 16), 598 (12), 596 (16), 500 (16), 499 (19), 498

(71), 497 (32), 496 (99), 173 (16), 116 (19), 57 (51); HRMS (FAB) calcd for

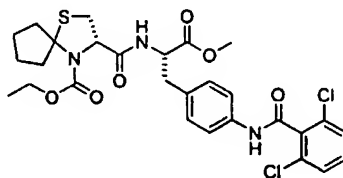
10 $\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}+\text{H}$, 596.1389, found 596.1364; Anal. Calcd for $\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.5$

H_2O : C, 53.56; H, 5.33; N, 6.94. Found: C, 53.86; H, 5.35; N, 6.90.

Example 77.

[S-(R*,R*)]-3-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-thia-4-azaspiro[4.4]nonane-4-carboxylic acid 4-ethyl ester

15 (Scheme B, B-6: where $\text{R}_{\text{B-1}}$ and $\text{R}_{\text{B-2}}$ are the same and equal to H, $\text{R}_{\text{B-3}}$ and $\text{R}_{\text{B-4}}$ together form a carbocyclic ring of 5 atoms, Y is CO_2 , R_3 is ethyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



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Example 77 was prepared as described in Scheme B from D-cysteine and cyclopentanone using ethyl chloroformate to form the requisite carbamate. Physical data as follows: IR

(mull) 1760, 1739, 1694, 1656, 1607, 1560, 1543, 1517, 1445, 1429, 1411, 1334, 1273,

1253, 1116 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.56 (2 H), 7.44 (1 H), 7.34 (3 H), 7.11 (2 H), 6.70

25 (1 H), 4.84 (2 H), 4.14 (2 H), 3.74 (3 H), 3.12 (4 H), 2.67 (1 H), 2.51 (1 H), 1.73 (6 H),

1.25 (3 H); ^{13}C NMR (CDCl_3) δ 171.4, 170.7, 162.3, 136.3, 135.8, 132.5, 132.4, 131.0,

130.1, 128.2, 120.2, 66.4, 62.1, 53.1, 52.5, 37.3, 32.3, 31.9, 25.1, 24.6, 14.5, 14.1; MS

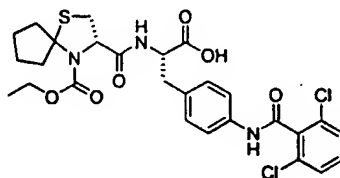
(ESI+) for $C_{28}H_{31}Cl_2N_3O_6S$ m/z 630.0 ($M+Na$)⁺; HRMS (EI) calcd for $C_{28}H_{31}Cl_2N_3O_6S$ 607.1310, found 607.1315; Anal. Calcd for $C_{28}H_{31}Cl_2N_3O_6S \cdot 0.75 H_2O$: C, 54.06; H, 5.27; N, 6.90. Found: C, 53.98; H, 5.16; N, 6.72.

Example 78.

5 [S-(R*,R*)]-3-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-1-thia-4-azaspiro[4.4]nonane-4-carboxylic acid 4-ethyl ester

(Scheme B, B-7: where $R_{B,1}$ and $R_{B,2}$ are the same and equal to H, $R_{B,3}$ and $R_{B,4}$ together form a carbocyclic ring of 5 atoms, Y is CO_2 , R_3 is ethyl, R_5 is 4-[(2,6-

10 dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



Example 78 was prepared from example 77 by the procedure described in preparation 6.

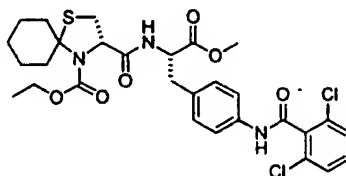
15 Physical data as follows: IR (mull) 3276, 1664, 1606, 1562, 1537, 1515, 1445, 1432, 1413, 1335, 1273, 1239, 1195, 1116, 799 cm^{-1} ; 1H NMR ($CDCl_3$) δ 8.08 (1 H), 7.62 (2 H), 7.24 (3 H), 7.14 (2 H), 6.72 (1 H), 4.85 (2 H), 4.13 (2 H), 3.21 (4 H), 2.53 (2 H), 1.75 (6 H), 1.22 (3 H); ^{13}C NMR ($CDCl_3$) δ 175.2, 171.2, 162.6, 136.7, 135.7, 132.3, 132.0, 130.7, 130.2, 127.9, 120.2, 66.3, 62.2, 53.2, 37.3, 36.6, 32.3, 25.2, 24.6, 20.5, 14.5; MS (ESI+) for $C_{27}H_{29}Cl_2N_3O_6S$ m/z 593.8 ($M+H$)⁺; MS (ESI-) for $C_{27}H_{29}Cl_2N_3O_6S$ m/z 591.8 ($M-H$)⁻; HRMS (FAB) calcd for $C_{27}H_{29}Cl_2N_3O_6S+H_1$ 594.1232, found 594.1226; Anal. Calcd for $C_{27}H_{29}Cl_2N_3O_6S \cdot 0.37 H_2O$: C, 53.95; H, 4.99; N, 6.99. Found: C, 54.28; H, 5.10; N, 7.03. % Water (KF): 1.10.

Example 79.

25 [S-(R*,R*)]-3-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-thia-4-azaspiro[4.5]decane-4-carboxylic acid 4-ethyl ester

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(Scheme B, B-6: where $R_{B,1}$ and $R_{B,2}$ are the same and equal to H, $R_{B,3}$ and $R_{B,4}$ together form a carbocyclic ring of 6 atoms, Y is CO_2 , R_3 is ethyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



5

Example 79 was prepared as described in Scheme B from D-cysteine and cyclohexanone using ethyl chloroformate to form the requisite carbamate. Physical data as follows: IR (mull) 1745, 1704, 1683, 1668, 1607, 1561, 1538, 1514, 1431, 1413, 1327, 1269, 1213, 1196, 1117 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.57 (2 H), 7.46 (1 H), 7.33 (3 H), 7.11 (2 H), 6.72 (1 H), 4.88 (2 H), 4.14 (2 H), 3.74 (3 H), 3.13 (4 H), 2.51 (1 H), 1.69 (8 H), 1.22 (5 H); ^{13}C NMR (CDCl_3) δ 171.4, 170.7, 168.1, 162.3, 136.3, 135.7, 132.4, 131.0, 128.2, 120.3, 66.9, 62.8, 62.0, 55.1, 53.1, 52.5, 42.0, 39.9, 37.3, 36.9, 31.1, 29.6, 27.6, 27.1, 26.0, 25.3, 24.7, 23.1, 14.5; MS (ESI-) for $\text{C}_{29}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 621.9 (M); HRMS (FAB) calcd for $\text{C}_{29}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_6\text{S} + \text{H}^+$ 622.1545, found 622.1536.

15

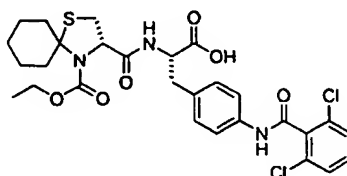
Example 80.

[S-(R*,R*)]-3-[[[1-Carboxy-2-[4-[(2,6-

dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-1-thia-4-azaspiro[4.5]decane-4-carboxylic acid 4-ethyl ester

(Scheme B, B-7: where $R_{B,1}$ and $R_{B,2}$ are the same and equal to H, $R_{B,3}$ and $R_{B,4}$ together form a carbocyclic ring of 6 atoms, Y is CO_2 , R_3 is ethyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).

20

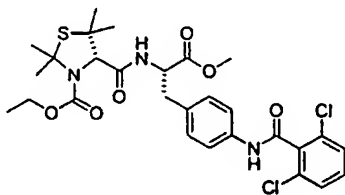


Example 80 was prepared from example 79 by the procedure described in preparation 6. Physical data as follows: IR (mull) 3276, 1710, 1664, 1606, 1562, 1537, 1515, 1432, 1413, 1329, 1272, 1256, 1195, 1117, 800 cm^{-1} ; ^1H NMR (CDCl_3) δ 8.07 (1 H), 7.62 (2 H), 7.24 (3 H), 7.14 (2 H), 6.75 (1 H), 4.88 (2 H), 4.12 (3 H), 3.14 (4 H), 2.74 (1 H), 2.50 (1 H), 1.69 (6 H), 1.19 (5 H); ^{13}C NMR (CDCl_3) δ 175.4, 175.1, 171.3, 162.6, 136.7, 135.7, 132.3, 132.0, 130.7, 130.3, 130.1, 128.0, 120.2, 66.8, 62.1, 53.2, 42.0, 36.9, 30.5, 27.0, 26.0, 25.3, 25.0, 24.6, 20.5, 14.4, 3.7; MS (ESI+) for $\text{C}_{28}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 608.1 ($\text{M}+\text{H}^+$); MS (ESI-) for $\text{C}_{28}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 605.9 ($\text{M}-\text{H}^-$); HRMS (EI) calcd for $\text{C}_{28}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ 607.1310, found 607.1309; Anal. Calcd for $\text{C}_{28}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.3 \text{H}_2\text{O}$: C, 54.78; H, 5.19; N, 6.85. Found: C, 54.56; H, 5.24; N, 6.90. % Water (KF): 0.87.

Example 81.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-2,2,5,5-tetramethyl-3-thiazolidinecarboxylic acid 3-ethyl ester

(Scheme B, B-6: where $\text{R}_{\text{B-1}}$, $\text{R}_{\text{B-2}}$, $\text{R}_{\text{B-3}}$ and $\text{R}_{\text{B-4}}$ are the same and equal to CH_3 , Y is CO_2 , R_3 is *t*-butyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



20

Example 81 was prepared as described in Scheme B from D-penicillamine and acetone using ethyl chloroformate to form the requisite carbamate. Physical data as follows: IR (mull) 1748, 1666, 1606, 1562, 1538, 1516, 1432, 1413, 1327, 1275, 1233, 1215, 1195, 1080, 799 cm^{-1} ; ^1H NMR (CDCl_3) δ 7.55 (2 H), 7.33 (4 H), 7.09 (3 H), 6.59 (1 H), 4.91 (1 H), 4.47 (1 H), 4.16 (2 H), 3.75 (3 H), 3.13 (2 H), 1.92 (3 H), 1.75 (3 H), 1.67 (3 H), 1.60 (3 H), 1.21 (3 H); ^{13}C NMR (CDCl_3) δ 171.5, 170.0, 162.2, 136.1, 135.7, 132.5, 132.3, 131.9, 130.9, 130.7, 129.7, 128.1, 120.3, 120.1, 120.0, 61.6, 61.4, 52.9, 52.4, 52.2,

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49.3, 37.4, 37.3, 34.1, 31.4, 30.8, 29.5, 24.9; MS (ESI-) for $C_{28}H_{33}Cl_2N_3O_6S$ m/z 607.9 (M-H); HRMS (EI) calcd for $C_{28}H_{33}Cl_2N_3O_6S$ 609.1467, found 609.1461; Anal. Calcd for $C_{28}H_{33}Cl_2N_3O_6S \cdot 0.19 H_2O$: C, 54.77; H, 5.48; N, 6.84. Found: C, 55.00; H, 5.48; N, 6.78. % Water (KF): 0.56.

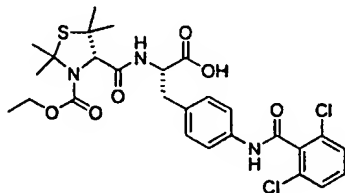
5

Example 82.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-2,2,5,5-tetramethyl-3-thiazolidinecarboxylic acid 3-ethyl ester

(Scheme B, B-7: where R_{B-1} , R_{B-2} , R_{B-3} and R_{B-4} are the same and equal to CH_3 , Y is CO_2 ,

10 R_1 is *t*-butyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



Example 82 was prepared from example 81 by the procedure described in preparation 6.

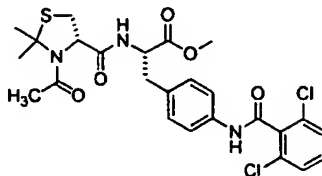
15 Physical data as follows: IR (mull) 3275, 1750, 1735, 1678, 1666, 1609, 1562, 1543, 1516, 1432, 1413, 1333, 1276, 1195, 1077 cm^{-1} ; 1H NMR (DMSO- d_6) δ 12.48 (1 H), 10.63 (1 H), 8.36 (1 H), 7.53 (5 H), 7.19 (2 H), 4.49 (2 H), 3.96 (2 H), 3.03 (1 H), 2.79 (1 H), 1.78 (6 H), 1.48 (3 H), 1.15 (3 H), 0.90 (3 H); ^{13}C NMR (DMSO- d_6) δ 172.9, 168.7, 161.7, 152.3, 137.0, 136.3, 132.9, 131.2, 131.1, 129.5, 128.1, 119.2, 74.7, 71.3, 60.5, 53.5, 49.0, 48.4, 38.3, 36.7, 33.5, 31.8, 28.1, 24.7, 21.0, 14.0; MS (ESI+) for $C_{27}H_{31}Cl_2N_3O_6S$ m/z 595.8 (M+H)⁺; MS (ESI-) for $C_{27}H_{31}Cl_2N_3O_6S$ m/z 593.8 (M-H)⁻; HRMS (FAB) calcd for $C_{27}H_{31}Cl_2N_3O_6S + H$, 596.1389, found 596.1362; Anal. Calcd for $C_{27}H_{31}Cl_2N_3O_6S \cdot 0.56 H_2O$: C, 53.46; H, 5.34; N, 6.93. Found: C, 53.73; H, 5.35; N, 6.73. % Water (KF): 1.67.

25 Example 83.

N-[[[(4S)-3-Acetyl-2,2-dimethyl-4-thiazolidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine methyl ester

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(Scheme B, B-6: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} are the same and equal to CH_3 , Y is CO, R_3 is methyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



5

Example 83 was prepared as described in Scheme B from D-cysteine and acetone using acetyl chloride to form the requisite amide. Physical data as follows: IR (mull) 1745, 1682, 1662, 1628, 1610, 1579, 1561, 1541, 1515, 1431, 1412, 1326, 1270, 1240, 1211 cm^{-1} ; 1H NMR ($CDCl_3$) δ 7.55 (3 H), 7.32 (3 H), 7.19 (1 H), 7.11 (1 H), 6.66 (1 H), 4.89 (1 H), 4.60 (1 H), 3.78 (3 H), 3.24 (4 H), 2.04 (3 H), 1.87 (3 H), 1.79 (3 H); ^{13}C NMR ($DMSO-d_6$) δ 171.6, 171.5, 169.9, 169.4, 167.9, 161.8, 137.1, 136.3, 132.8, 132.6, 131.1, 129.5, 128.1, 119.2, 72.6, 66.5, 53.7, 53.3, 52.0, 51.9, 35.8, 31.7, 29.0, 26.9, 24.7, 24.5; MS (ESI+) for $C_{25}H_{27}Cl_2N_3O_5S$ m/z 551.9 ($M+H$) $^+$; HRMS (EI) calcd for $C_{25}H_{27}Cl_2N_3O_5S$ 551.1049, found 551.1053; MS (EI) m/z (rel. intensity) 551 (M^+ , 7), 351 (46), 349 (68), 278 (16), 186 (14), 175 (63), 173 (98), 158 (23), 116 (99), 100 (20), 99 (69).

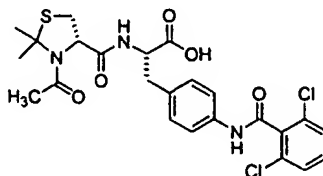
15

Example 84.

N-[[[(4S)-3-Acetyl-2,2-dimethyl-4-thiazolidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine

(Scheme B, B-7: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} are the same and equal to CH_3 , Y is CO, R_3 is methyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).

20



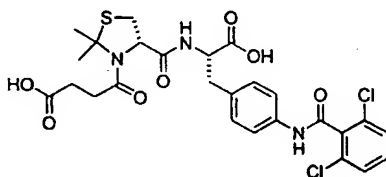
Example 84 was prepared from example 83 by the procedure described in preparation 6.

Physical data as follows: IR (mull) 3279, 1723, 1661, 1608, 1562, 1542, 1516, 1432, 1413, 1349, 1329, 1270, 1238, 1207, 1195 cm^{-1} ; ^1H NMR (DMSO-d_6) δ 12.47 (1 H), 10.62 (1 H), 7.51 (5 H), 7.18 (2 H), 4.80 (1 H), 4.67 (1 H), 4.47 (1 H), 2.98 (3 H), 1.68 (9 H); ^{13}C NMR (DMSO-d_6) δ 184.2, 172.6, 171.9, 169.8, 169.2, 167.8, 136.9, 136.3, 133.1, 131.2, 131.0, 129.5, 128.1, 119.2, 72.6, 66.6, 53.7, 53.3, 36.2, 31.7, 29.0, 27.0, 24.8, 24.6, 21.0; MS (ESI+) for $\text{C}_{24}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_5\text{S}$ m/z 538.0 ($\text{M}+\text{H}^+$); MS (ESI-) for $\text{C}_{24}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_5\text{S}$ m/z 535.9 ($\text{M}-\text{H}^-$); HRMS (FAB) calcd for $\text{C}_{24}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_5\text{S}+\text{H}$, 538.0970, found 538.0961.

Example 85.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-2,2-dimethyl- γ -oxo-3-thiazolidinebutanoic acid

(Scheme B, B-7: where $\text{R}_{\text{B-1}}$ and $\text{R}_{\text{B-2}}$ are the same and equal to H, $\text{R}_{\text{B-3}}$ and $\text{R}_{\text{B-4}}$ are the same and equal to CH_3 , Y is CO, R_3 is $\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



- Example 85 was prepared as described in Scheme B from D-cysteine and acetone using methyl succinyl chloride to form the requisite amide. Physical data as follows: IR (mull) 3264, 3125, 3071, 1724, 1658, 1607, 1562, 1537, 1517, 1432, 1414, 1326, 1241, 1195, 1181 cm^{-1} ; ^1H NMR (DMSO-d_6) δ 12.31 (1 H), 10.65 (1 H), 8.20 (1 H), 7.50 (5 H), 7.19 (2 H), 5.81 (1 H), 4.50 (1 H), 3.02 (3 H), 2.30 (3 H), 1.71 (6 H); ^{13}C NMR (DMSO-d_6) δ 174.1, 172.8, 169.4, 161.9, 137.1, 136.5, 133.2, 131.4, 131.3, 129.7, 128.3, 119.3, 73.1, 65.7, 53.5, 36.3, 32.0, 30.5, 29.1, 29.0, 27.1; MS (ESI+) for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 595.9 ($\text{M}+\text{H}^+$); MS (ESI-) for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 593.8 ($\text{M}-\text{H}^-$); Anal. Calcd for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$

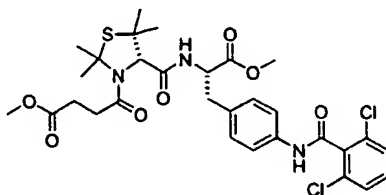
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• 0.51 H₂O: C, 51.55; H, 4.66; N, 6.94. Found: C, 51.71; H, 4.85; N, 6.93. % Water (KF): 1.53.

Example 86.

5 [S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-2,2,5,5-tetramethyl-γ-oxo-3-thiazolidinebutanoic acid methyl ester

(Scheme B, B-6: where R_{B-1}, R_{B-2}, R_{B-3} and R_{B-4} are the same and equal to CH₃, Y is CO, R₃ is CH₂CH₂CO₂CH₃, R₄ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is
10 (S, S)).

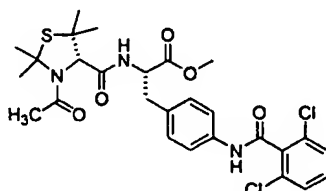


Example 86 was prepared as described in Scheme B from D-penicillamine and acetone
15 using methyl succinyl chloride to form the requisite amide. Physical data as follows: IR (mull) 3287, 1741, 1660, 1608, 1562, 1540, 1516, 1432, 1413, 1323, 1267, 1241, 1225, 1196, 1168 cm⁻¹; ¹H NMR (CDCl₃) δ 7.55 (2 H), 7.29 (3 H), 7.13 (2 H), 6.94 (1 H), 4.95 (1 H), 4.39 (1 H), 3.74 (3 H), 3.67 (3 H), 3.27 (1 H), 3.10 (1 H), 2.80 (1 H), 2.57 (3 H), 2.18 (1 H), 1.95 (3 H), 1.77 (3 H), 1.72 (3 H), 1.68 (3 H) 1.63 (3 H); ¹³C NMR (CDCl₃) δ
20 173.3, 171.6, 170.3, 169.9, 162.4, 136.3, 132.5, 132.4, 131.0, 130.2, 129.8, 128.2, 120.8, 120.7, 120.5, 73.8, 53.1, 52.6, 51.8, 49.9, 40.0, 37.4, 33.9, 31.6, 31.4, 29.4, 29.1, 24.3; MS (ESI+) for C₃₀H₃₅Cl₂N₃O₇S *m/z* 652.1 (M+H)⁺; Anal. Calcd for C₃₀H₃₅Cl₂N₃O₇S • 0.31 H₂O: C, 54.75; H, 5.45; N, 6.38. Found: C, 55.04; H, 5.50; N, 6.69. % Water (KF): 0.84.

Example 87.

25 N-[[[(4S)-3-Acetyl-2,2,5,5-tetramethyl-4-thiazolidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine methyl ester

(Scheme B, B-6: where R_{B-1} , R_{B-2} , R_{B-3} and R_{B-4} are the same and equal to CH_3 , Y is CO, R_3 is methyl, R_4 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



5

Example 87 was prepared as described in Scheme B from D-penicillamine and acetone using acetyl chloride to form the requisite amide. Physical data as follows: IR (mull) 1741, 1692, 1676, 1629, 1606, 1563, 1538, 1516, 1431, 1410, 1352, 1320, 1276, 1251, 1223 cm^{-1} ; 1H NMR (DMSO- d_6) δ 10.70 (1 H), 8.57 (1 H), 7.56 (4 H), 7.22 (2 H), 4.63 (1 H), 4.49 (1 H), 3.65 (3 H), 3.30 (1 H), 3.12 (1 H), 2.84 (1 H), 1.92 (3 H), 1.82 (3 H), 1.78 (3 H), 1.51 (3 H), 0.80 (3 H); ^{13}C NMR (DMSO- d_6) δ 184.2, 171.9, 168.9, 168.1, 161.8, 137.1, 136.3, 132.4, 131.3, 131.2, 131.0, 129.4, 128.1, 119.3, 75.9, 72.8, 53.5, 52.0, 49.2, 48.4, 36.0, 33.5, 31.9, 27.6, 24.9, 24.5; MS (ESI+) for $C_{27}H_{31}Cl_2N_3O_5S$ m/z 580.1 (M+H) $^+$; MS (ESI-) for $C_{27}H_{31}Cl_2N_3O_5S$ m/z 577.9 (M-H) $^-$; Anal. Calcd for $C_{27}H_{31}Cl_2N_3O_5S$ • 0.12 H_2O : C, 55.66; H, 5.40; N, 7.21. Found: C, 55.68; H, 5.39; N, 7.16. % Water (KF): 0.36.

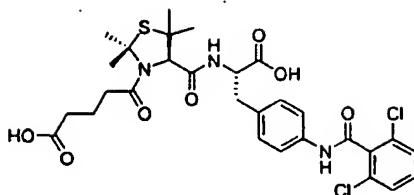
20

Example 88.

[R-(R*,S*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-2,2,5,5-tetramethyl- δ -oxo-3-thiazolidinepentanoic acid

(Scheme B, B-6: where R_{B-1} , R_{B-2} , R_{B-3} and R_{B-4} are the same and equal to CH_3 , Y is CO, R_3 is $CH_2CH_2CH_2CO_2H$, R_4 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (R, S)).

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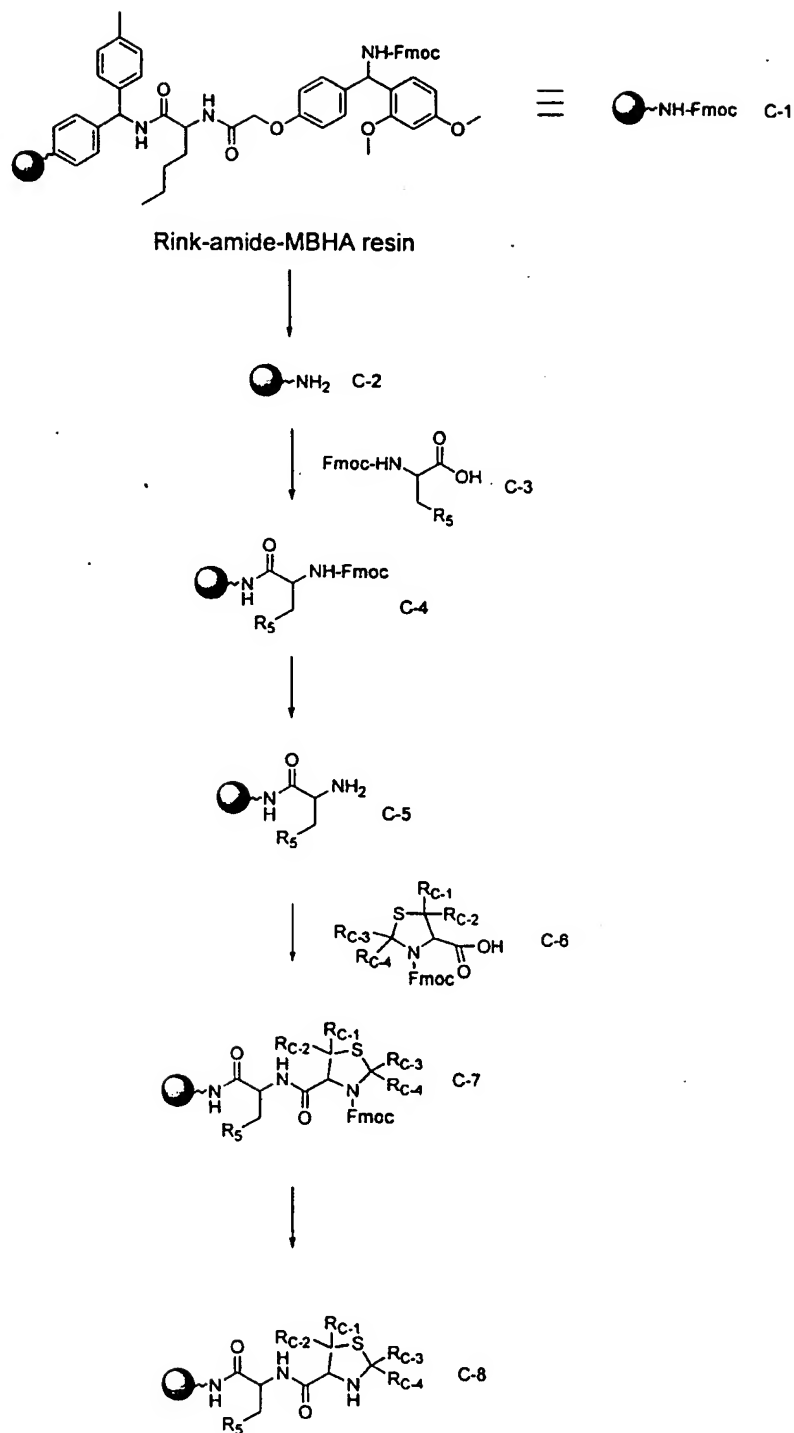


Example 88 was prepared as described in Scheme B from D-penicillamine and acetone using methyl glutaryl chloride to form the requisite amide. Physical data as follows:

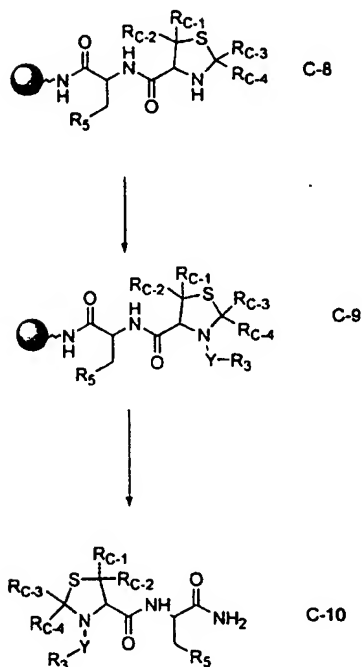
- 5 ^1H NMR (DMSO- d_6) δ 10.65 (1 H), 8.45 (1 H), 7.53 (5 H), 7.22 (2 H), 4.51 (1 H), 3.11 (3 H), 2.77 (1 H), 2.21 (4 H), 1.89 (3 H), 1.82 (3 H), 1.11 (2 H), 1.48 (3 H), 0.77 (3 H); ^{13}C NMR (DMSO- d_6) δ 184.2, 174.2, 173.1, 170.1, 168.7, 161.7, 137.0, 136.3, 133.1, 131.3, 131.1, 129.4, 128.1, 119.2, 74.9, 73.0, 53.8, 49.3, 36.1, 34.8, 33.5, 32.7, 31.9, 27.7, 24.5, 20.0; MS (ESI+) for $\text{C}_{29}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 638.0 (M+H) $^+$; MS (ESI-) for
- 10 $\text{C}_{29}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_7\text{S}$ m/z 635.9 (M-H) $^-$; HRMS (FAB) calcd for $\text{C}_{29}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_7\text{S} + \text{H}_1$ 638.1494, found 638.1481; Anal. Calcd for $\text{C}_{29}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_7\text{S} \cdot 0.75 \text{H}_2\text{O}$: C, 53.42; H, 5.33; N, 6.44. Found: C, 53.20; H, 5.26; N, 6.45.

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Scheme C



Scheme C (continued)



5 Where: R_{C-1}, R_{C-2}, and R_{C-3} are defined independently as R₁. R_{C-4} is defined as R₂.

Scheme C describes a method for the preparation of examples of the formula C-10. Commercially available Rink Amide MBHA resin is deprotected under standard solid-phase peptide synthesis conditions (Atherton, E.; Sheppard R.C. *Solid Phase Peptide Synthesis: A Practical Approach*; IRL Press at Oxford University Press: Oxford, 1989) to afford the amine of formula C-2. Acylation with a commercially available or readily prepared amino acid residue of general C-3 affords the resin bound derivative of formula C-4. Removal of the Fmoc group under standard conditions provides amine of general structure C-5 which is acylated with a commercially available or readily prepared thiazolidine-4-carboxylic acid of general formula C-6 to afford the resin bound intermediate C-7. Standard Fmoc deprotection affords the resin bound amine of general formula C-8 which may be reacted with a variety of electrophilic reagents as described in Scheme A to afford resin bound amides, ureas, sulfonamides and carbamates of general structure C-9. Preparation 18 details an example of the reaction of a mixed carbonate to

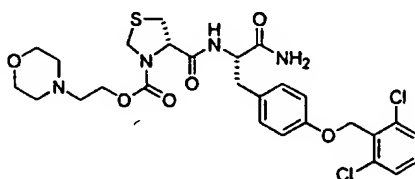
afford a carbamate of general structure C-9 (where Y is equal to CO₂). Standard acidolysis affords the amide of general structure C-10.

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Preparation 18 and Example 89.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-amino-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(4-morpholinyl)ethyl] ester

(Scheme C, C-10: where R_{C-1}, R_{C-2}, R_{C-3} and R_{C-4} are the same and equal to proton, R₃ is 2-(4-morpholinyl)ethyl, R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl, Y is CO₂ and stereochemistry is (S, S)).



15

To a mixture of Rink Amide MBHA resin (Scheme C, C-1) (Nova Biochem., 1.2 g, ca. 0.59 mmol) in methylene chloride (20 mL) was added a solution of piperidine in DMF (30%, 20 mL). A slow stream of nitrogen was bubbled through the mixture to effect mixing for 20 min. The resin was filtered and washed with DMF. The resin was suspended in a solution of piperidine in DMF (30%, 40 ml) and mixed for 40 min. The resin was filtered and washed with DMF, methylene chloride, methanol and methylene chloride to afford resin C-2, which was diluted with DMF (40 mL). To this mixture was added Fmoc-Tyr(2,6-Cl₂-Bn) (Scheme C, C-3: where R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S)) (Advanced Chemtech, 1.32 g, 2.35 mmol), HOBt (0.36 g, 2.35 mmol), PyBOP (1.20 g, 2.35) and DIEA (1.03 mL, 5.90 mmol). The reaction was mixed for 4 h and the resin filtered and washed with DMF, methylene chloride, MeOH and methylene chloride to afford the intermediate resin-bound

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amino acid derivative C-4 (Scheme C, where R_3 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*)) which was used without characterization.

To a mixture of resin C-4 described above in methylene chloride (20 mL) was added a solution of piperidine in DMF (30%, 20 mL). A slow stream of nitrogen was bubbled through the mixture to effect mixing for 20 min. The resin was filtered and washed with DMF. The resin was suspended in a solution of piperidine in DMF (30%, 40 ml) and mixed for 40 min. The resin was filtered and washed with DMF, methylene chloride, methanol and methylene chloride to afford the resin of structure C-5 (Scheme C, where R_3 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*)), which was diluted with DMF (40 mL). To this mixture was added Fmoc-D-thiazolidine-4-carboxylic acid (Scheme C, C-6: where R_{C-1} , R_{C-2} , R_{C-3} and R_{C-4} are the same and equal to proton and stereochemistry is (*S*)) (Advanced Chemtech, 832 mg, 2.35 mmol), HOBt (0.36 g, 2.35 mmol), PyBOP (1.20 g, 2.35) and DIEA (1.03 mL, 5.90 mmol). The reaction was mixed for 4 h and the resin filtered and washed with DMF, methylene chloride, MeOH and methylene chloride to afford the intermediate resin-bound derivative C-7 (Scheme C, where R_{C-1} , R_{C-2} , R_{C-3} and R_{C-4} are the same and equal to proton, R_3 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*, *S*)) which was used without characterization.

To a mixture of resin C-7 described above in methylene chloride (20 mL) was added a solution of piperidine in DMF (30%, 20 mL). A slow stream of nitrogen was bubbled through the mixture to effect mixing for 20 min. The resin was filtered and washed with DMF. The resin was suspended in a solution of piperidine in DMF (30%, 40 ml) and mixed for 40 min. The resin was filtered and washed with DMF, methylene chloride, methanol and methylene chloride to provide the intermediate of structure C-8 (Scheme C, where R_{C-1} , R_{C-2} , R_{C-3} and R_{C-4} are the same and equal to proton, R_3 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*, *S*)), which was diluted with methylene chloride (10 mL). To this mixture was added a solution of the mixed carbonate prepared from 4-(2-hydroxyethyl)morpholine (2.14 mL, 17.7 mmol) and N,N-disuccinimidyl carbonate (4.53 g, 17.7 mmol) as described in preparation 8 in methylene chloride (20 mL) followed by triethylamine (0.33 mL, 2.36 mmol). The reaction was

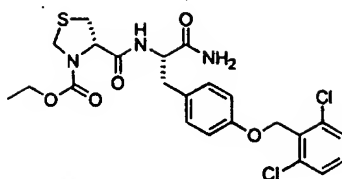
mixed for 24 h and the resin filtered and washed extensively with DMF, methylene chloride, MeOH, and anhydrous ethyl ether. The resin was dried *in vacuo* to afford the resin-bound carbamate C-9 (Scheme C, where R_{C-1} , R_{C-2} , R_{C-3} and R_{C-4} are the same and equal to proton, R_3 is 2-(4-morpholinyl)ethyl, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, Y is CO_2 and stereochemistry is (S, S)).

Resin C-9 was swelled with a minimum of methylene chloride (ca. 2 mL) and suspended with 95 % aqueous TFA (20 mL). The mixture was mixed by magnetic stirring for 1h and filtered. The resin was washed with additional TFA (2 X 5 mL), followed by methylene chloride and methanol. The combined filtrates were evaporated *in vacuo*, and partitioned between ethyl acetate and saturated aqueous sodium bicarbonate. The organic layer was separated and washed with brine, dried ($MgSO_4$), filtered and evaporated *in vacuo*. The residue was purified by flash chromatography using methylene chloride/methanol (1 to 3%) as eluant to afford the title compound (215 mg) as an amorphous powder: IR (mull) 3288, 1676, 1657, 1611, 1564, 1511, 1439, 1424, 1346, 1302, 1237, 1179, 1116, 1021, 767 cm^{-1} ; 1H NMR ($CDCl_3$, 300 MHz) δ 7.24 (5 H), 6.93 (2 H), 5.20 (2 H), 4.63 (3 H), 4.34 (1 H), 4.22 (2 H), 3.63 (4 H), 3.11 (4 H), 2.50 (6 H); ^{13}C NMR ($CDCl_3$, 75 MHz) δ 173.4, 170.1, 157.9, 154.5, 136.9, 132.0, 130.4, 129.1, 128.4, 115.1, 66.8, 65.2, 63.4, 57.1, 54.0, 53.7, 53.5, 49.4, 35.2; MS (EI) m/z (rel. intensity) 610 (M^+ , 1), 323 (13), 321 (20), 161 (34), 159 (53), 114 (31), 113 (98), 100 (99), 88 (13), 70 (8), 56 (11); MS (FAB) m/z (rel. intensity) 611 (MH^+ , 71), 614 (18), 613 (49), 612 (27), 611 (71), 123 (60), 114 (99), 113 (76), 112 (19), 107 (22), 100 (28); HRMS (FAB) calcd for $C_{27}H_{32}Cl_2N_4O_6S + H$, 611.1498, found 611.1494. Anal. Calcd for $C_{27}H_{32}Cl_2N_4O_6S$: C, 53.03; H, 5.27; N, 9.16. Found: C, 52.74; H, 5.17; N, 9.01.

Example 90.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-amino-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester (Scheme C, C-10: where R_{C-1} , R_{C-2} , R_{C-3} and R_{C-4} are the same and equal to proton, R_3 is ethyl, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, Y is CO_2 and stereochemistry is (S, S)).

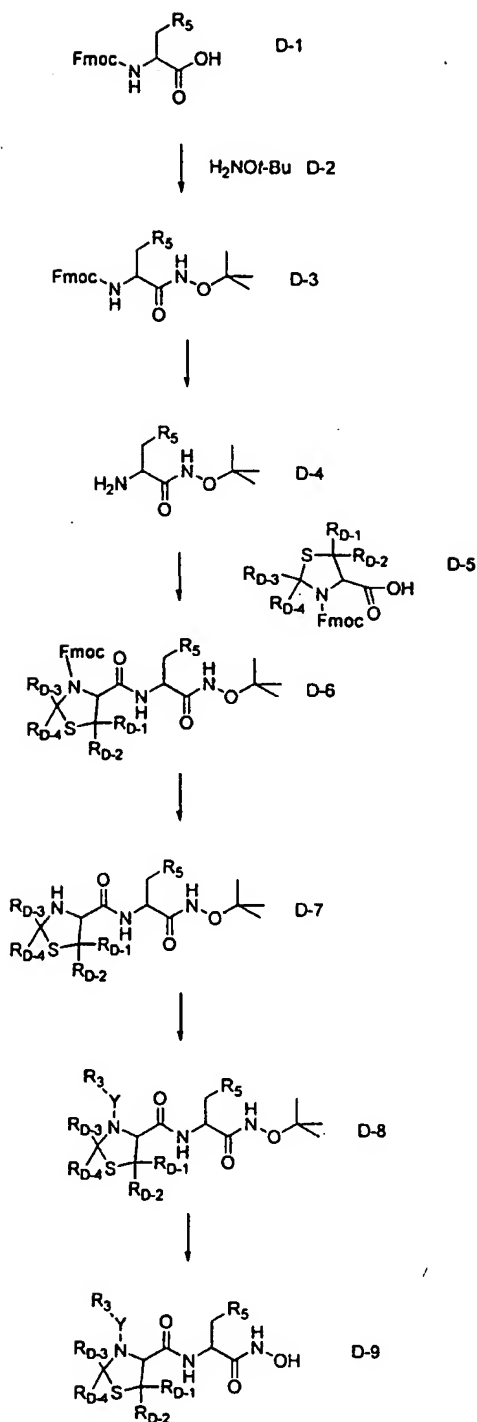
-111-



Example 90 was prepared as described in Scheme C. Physical data as follows: IR (mull) 3369, 3308, 3192, 1713, 1667, 1650, 1629, 1539, 1513, 1441, 1344, 1290, 1240, 1016, 768 cm^{-1} ; ^1H NMR (CDCl_3 , 300 MHz) δ 7.33 (3 H), 7.16 (2 H), 6.95 (2 H), 5.25 (2 H), 4.48 (4 H), 4.20 (2 H), 3.03 (4 H), 1.26 (3 H); ^{13}C NMR (CDCl_3 , 75 MHz) δ 173.6, 170.5, 157.8, 154.7, 136.8, 313.9, 130.4, 130.2, 129.0, 128.4, 114.9, 67.9, 65.1, 63.1, 62.5, 53.8, 36.5, 14.3; MS (EI) m/z (rel. intensity) 525 (M^+ , 1), 323 (44), 322 (13), 321 (68), 267 (9), 265 (14), 163 (12), 161 (65), 160 (35), 159 (99), 88 (30); MS (FAB) m/z (rel. intensity) 526 (MH^+ , 58), 528 (40), 527 (19), 526 (58), 321 (27), 188 (29), 161 (37), 160 (99), 159 (48), 107 (26), 88 (39); HRMS (FAB) calcd for $\text{C}_{23}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_5\text{S} + \text{H}$, 526.0970, found 526.0942. Anal. Calcd for $\text{C}_{23}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_5\text{S}$: C, 52.47; H, 4.79; N, 7.98. Found: C, 52.34; H, 4.81; N, 7.90.

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Scheme D

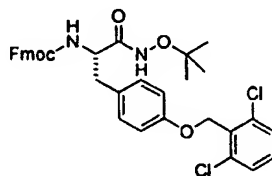


R_{D-1} , R_{D-2} , and R_{D-3} are defined independently as R_1 . R_{D-4} is defined as R_2 .

Scheme D describes a method for the preparation of examples of general formula **D-9**. Commercially available or readily prepared N- α -Fmoc protected amino acids of general structure **D-1** are coupled with *O*-(*tert*-butyl)hydroxylamine (**D-2**) under standard
5 coupling conditions as previously referenced to afford the *t*-butyl hydroxamate of general structure **D-3**. Standard Fmoc deprotection affords the intermediate amine of formula **D-4**. Coupling of this amine with a commercially available or readily prepared N- α -Fmoc-thiazolidine-4-carboxylic acid of general structure **D-5** affords the pseudodipeptide intermediate of general structure **D-6**. Standard Fmoc deprotection affords the
10 intermediate amine of general structure **D-7** which may be reacted under the variety of conditions described in Scheme A to afford amides, carbamates, sulfonamides and ureas of general structure **D-8**. Preparation 22 provides a specific example of the reaction of an amine of general structure **D-7** with a carbonate to afford a carbamate of general structure **D-8**. Mild acidolysis affords the hydroxamate of general structure **D-9**.

15 Preparation 19.

(Scheme D, **D-3**: where R_2 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*)).

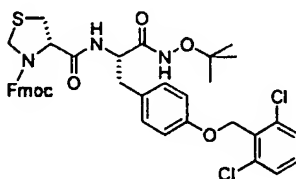


20 To a cooled (0-5 °C) solution of Fmoc-Tyr(2,6-Cl₂-Bn) (Scheme D, **D-1**: where R_2 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*)) (Advanced Chemtech, 6.0 g, 10.7 mmol), HOBt (1.63 g, 10.7 mmol), *O*-(*tert*-butyl)hydroxylamine hydrochloride (Scheme D, **D-2**) (1.61 g, 12.80 mmol) in methylene chloride (30 mL) was added PyBOP (6.66 g, 12.80 mmol) followed by DIEA (6.51 mL, 37.35 mmol). The mixture was stirred
25 at 0-5 °C for 1 hour, gradually allowed to warm to room temperature and stirred an additional 2 h. The mixture was diluted with methylene chloride and 0.25 N HCl, the organic layer separated and washed with saturated aqueous NaHCO₃, and brine, dried

(MgSO₄), filtered and evaporated *in vacuo*. The residue was purified by flash chromatography using methylene chloride/methanol (0-2.5%) as eluant to afford the title compound (5.87 g) as an amorphous powder: ¹H NMR (CDCl₃, 300 MHz) δ 8.08 (1 H), 7.76 (2 H), 7.54 (2 H), 7.29 (9 H), 6.95 (2 H), 5.43 (1 H), 5.30 (2 H), 4.28 (4 H), 3.06 (2 H), 1.19 (9 H); ¹³C NMR (CDCl₃, 75 MHz) δ 169.7, 157.8, 156.3, 143.6, 141.2, 136.8, 132.0, 130.4, 128.9, 128.4, 127.7, 127.0, 125.0, 119.9, 115.0, 82.3, 67.1, 65.1, 54.0, 46.9, 37.6, 26.0; MS (ESI+) for C₃₅H₃₄Cl₂N₂O₅, *m/z* 632.9 (M+H)⁺. MS (ESI+) for C₃₅H₃₄Cl₂N₂O₅, *m/z* 654.9 (M+Na)⁺.

Preparation 20.

- 10 (Scheme D, **D-6**: where R_{D-1}, R_{D-2}, R_{D-3} and R_{D-4} are the same and equal to proton, R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*, *S*)).



- 15 To a solution of **D-3** (Scheme D, where R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*)) (5.87 g, 9.28 mmol) in anhydrous DMF (94 mL) was added diethylamine (9.40 mL, 90.84 mmol) at ambient temperature. The solution was stirred for 90 min and volatiles were removed *in vacuo* to afford the intermediate amine **D-4** (Scheme D, where R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*)) as an oil which was used without further purification.
- 20

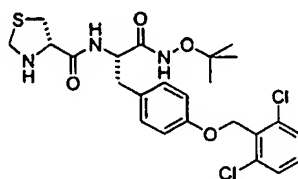
- To a cooled (0-5 °C) solution of Fmoc-D-thiazolidine-4-carboxylic acid (Scheme D, **D-5**: where R_{D-1}, R_{D-2}, R_{D-3} and R_{D-4} are the same and equal to proton and stereochemistry is (*S*)) (Advanced Chemtech, 3.93 g, 11.10 mmol) and HOAt (1.51 g, 11.10 mmol) in methylene chloride/DMF (4:1, 30 mL) was added EDC (2.12 g, 11.10 mmol). The reaction mixture was allowed to stir for 15 min and a solution of the amine (**D-4** described above) in methylene chloride/DMF (4:1, 30 mL) was added followed by
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DIEA (1.61 mL, 9.28 mmol). After 1 h at 0-5 °C, an additional equivalent of DIEA (1.61 mL, 9.28 mmol) was added and the mixture allowed to warm to room temperature. After stirring overnight, volatiles were removed *in vacuo* and the residue partitioned between ethyl acetate and 0.25 N aqueous HCl. The organic layer was separated and washed with water, saturated aqueous NaHCO₃, brine, dried (Na₂SO₄), filtered and concentrated *in vacuo*. The residue was purified by flash chromatography using CH₂Cl₂/acetone (3%) containing isopropanol (0.1%) as eluant to afford the title compound (2.4 g) as an amorphous solid: ¹H NMR (300 MHz, CDCl₃) δ 7.77 (2 H), 7.55 (2 H), 7.32 (7 H), 7.12 (2 H), 6.92 (2 H), 6.70 (1 H), 5.19 (3 H), 4.55 (5 H), 4.26 (2 H), 3.30 (1 H), 3.11 (3 H), 1.14 (9 H); ¹³C NMR (75 MHz, CDCl₃) δ 170.0, 168.8, 158.0, 143.4, 141.3, 137.0, 132.1, 130.4, 128.5, 127.9, 127.2, 124.9, 120.1, 115.3, 82.5, 68.4, 65.3, 52.8, 47.1, 36.5, 26.2. MS (ESI+) for C₃₉H₃₉Cl₂N₃O₆S *m/z* 747.9 (M+H)⁺; MS (ESI+) for C₃₉H₃₉Cl₂N₃O₆S *m/z* 769.8 (M+Na)⁺; MS (ESI-) for C₃₉H₃₉Cl₂N₃O₆S *m/z* 745.7 (M-H)⁻.

Preparation 21.

(Scheme D, D-7: where R_{D-1}, R_{D-2}, R_{D-3} and R_{D-4} are the same and equal to proton, R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).

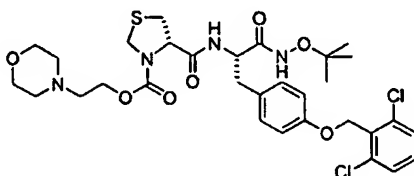


To a solution of D-6 (Scheme D, where R_{D-1}, R_{D-2}, R_{D-3} and R_{D-4} are the same and equal to proton, R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)) (500 mg, 0.67 mmol) in anhydrous DMF (7 mL) was added diethylamine (0.70 mL, 6.55 mmol) at ambient temperature. The solution was stirred for 90 min and volatiles were removed *in vacuo*. The residue was washed with ethyl ether/hexane (3:2) to afford the title compound (352 mg) as an amorphous solid which was used without further purification: ¹H NMR (300 MHz, DMSO-d₆) δ 10.61 (1 H), 8.27 (1 H), 7.54 (2 H), 7.44 (1 H), 7.15 (2 H), 6.94 (2 H), 5.15 (2 H), 4.50 (1 H), 4.03 (2 H), 3.75 (1 H), 3.19 (1 H), 2.82 (3 H), 2.57 (1 H),

1.06 (9 H); MS (ESI+) for $C_{24}H_{29}Cl_2N_3O_4S$ m/z 526.1 (M+H)⁺; MS (ESI-) for $C_{24}H_{29}Cl_2N_3O_4S$ m/z 524.1 (M-H)⁻.

Preparation 22.

(Scheme D, **D-8**: where R_{D-1} , R_{D-2} , R_{D-3} and R_{D-4} are the same and equal to proton, R_3 is 2-(4-morpholinyl)ethyl, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, Y is CO₂ and stereochemistry is (*S*, *S*)).



To a solution of 4-(2-hydroxyethyl)morpholine (1.22 mL, 10.05 mmol) in CH₃CN (55 mL) at ambient temperature was added N,N-disuccinimidyl carbonate (2.49 g, 10.05 mmol) and triethylamine (4.20 mL, 30.15 mmol). The solution was stirred at room temperature for 4 h and concentrated *in vacuo* to give a viscous oil. The oil was dissolved in a minimal amount of methylene chloride (15 mL) and added to a solution of **D-7** (Scheme D, where R_{D-1} , R_{D-2} , R_{D-3} and R_{D-4} are the same and equal to proton, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*, *S*) (350 mg, 0.67 mmol), triethylamine (0.10 mL, 0.74 mmol) and DMAP (1 mg) in CH₂Cl₂ (4 mL). The reaction mixture was stirred overnight and diluted with CH₂Cl₂ (15 mL). Propylamine (8.6 mL, 100.5 mmol) was slowly added (*exothermic*) and the solution stirred vigorously for 15 min, then diluted with water. The organic layer was separated and washed with 0.1 M HCl, saturated aqueous NaHCO₃, and brine, dried (MgSO₄), filtered and concentrated *in vacuo*. Purification of the residue by flash chromatography using ethyl acetate/acetone (3:1) as eluant afforded the title compound (251 mg) as a white powder: IR (mull) 3264, 1709, 1661, 1564, 1531, 1512, 1439, 1419, 1345, 1301, 1241, 1181, 1118, 1016, 767 cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 8.48 (1 H), 7.36 (2 H), 7.24 (1 H), 7.17 (2 H), 6.96 (2H), 5.23 (2 H), 4.60 (3 H), 4.31 (3 H), 3.71 (4 H), 3.33 (4 H), 2.59 (6 H), 1.15 (9 H); ¹³C NMR (75 MHz, CDCl₃) δ 168.9, 158.0, 137.0, 132.1, 130.5, 128.7, 128.5, 115.3, 82.5,

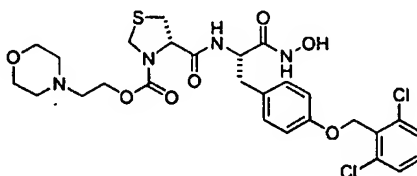
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66.8, 65.3, 63.4, 57.2, 53.7, 52.9, 36.4, 30.6, 29.3, 26.2, 19.1, 13.7; MS (ESI+) for $C_{31}H_{40}Cl_2N_4O_7S$ m/z 682.9 (M+H)⁺. MS (ESI+) for $C_{31}H_{40}Cl_2N_4O_7S$ m/z 705.0 (M+Na)⁺; Anal. Calcd for $C_{31}H_{40}Cl_2N_4O_7S \cdot 0.35 H_2O$: C, 53.97; H, 5.95; N, 7.95. Found: C, 54.22; H, 6.11; N, 7.95. % Water (KF): 0.91.

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Preparation 23 and Example 91.

(Scheme D, **D-9**: where R_{D-1} , R_{D-2} , R_{D-3} and R_{D-4} are the same and equal to proton, R_3 is 2-(4-morpholinyl)ethyl, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, Y is CO₂ and stereochemistry is (S, S)).



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Hydroxamate **D-8** (Scheme D, where R_{D-1} , R_{D-2} , R_{D-3} and R_{D-4} are the same and equal to proton, R_3 is 2-(4-morpholinyl)ethyl, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, Y is CO₂ and stereochemistry is (S, S)) (150 mg, 0.22 mmol) was dissolved in anhydrous TFA (12 mL) at ambient temperature and gradually warmed to 40 °C. After 5 h at 40 °C, volatiles were removed *in vacuo* and the residue partitioned between ethyl acetate and saturated aqueous NaHCO₃. The organic layer was separated and washed with brine, dried (MgSO₄), filtered and concentrated *in vacuo*. Purification of the residue by flash chromatography using methylene chloride/methanol (5%) as eluant afforded the title compound (51 mg) as an amorphous solid: IR (mull) 3273, 3229, 1708, 1652, 1564, 1546, 1511, 1439, 1422, 1346, 1236, 1180, 1114, 1022, 768 cm⁻¹; ¹H NMR (300 MHz, DMSO-d₆) δ 10.70 (1 H), 8.93 (1 H), 8.32 (1 H), 7.54 (2 H), 7.45 (1 H), 7.12 (2 H), 6.93 (2 H), 5.16 (2 H), 4.56 (2 H), 4.36 (1 H), 4.25 (1 H), 4.06 (2 H), 3.51 (4 H), 3.15 (1 H), 2.76 (3 H), 2.36 (4 H); ¹³C NMR (75 MHz, DMSO-d₆) δ 169.8, 167.9, 157.6, 136.5, 132.2, 132.0, 130.8, 130.5, 129.2, 114.7, 66.7, 65.3, 63.3, 59.1, 57.1, 53.8, 52.1, 37.8, 31.3, 30.1; MS (ESI+) for $C_{27}H_{32}Cl_2N_4O_7S$ m/z 627.0 (M+H)⁺; MS (ESI-) for $C_{27}H_{32}Cl_2N_4O_7S$ m/z 624.9

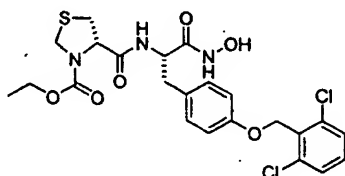
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-118-

(M-H); Anal. Calcd for $C_{27}H_{32}Cl_2N_4O_7S \cdot 0.46 H_2O$: C, 51.00; H, 5.22; N, 8.81. Found: C, 51.34; H, 5.23; N, 8.67. % Water (KF): 1.31.

Example 92.

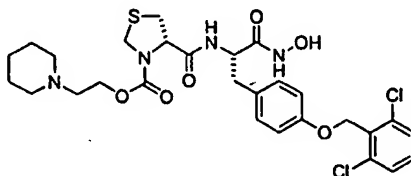
- (Scheme D, D-9: where R_{D-1} , R_{D-2} , R_{D-3} and R_{D-4} are the same and equal to proton, R_3 is ethyl, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, Y is CO_2 and stereochemistry is (S, S)).



- Example 92 was prepared as described in Scheme D from Fmoc-Tyr(2,6- Cl_2 -Bn) using ethyl chloroformate to provide the requisite carbamate. Physical properties as follows: IR (mull) 3278, 1654, 1612, 1585, 1564, 1547, 1511, 1439, 1347, 1237, 1195, 1179, 1022, 782, 769 cm^{-1} ; 1H NMR (300 MHz, $CDCl_3$) δ 7.39 (3 H), 7.17 (2 H), 6.94 (2 H), 5.24 (2 H), 4.53 (4 H), 4.18 (2 H), 3.15 (2 H), 2.87 (2 H), 1.28 (3 H); ^{13}C NMR (75 MHz, CD_3OD) δ 172.7, 170.1, 159.4, 156.2, 138.1, 133.7, 132.1, 131.6, 130.8, 129.8, 115.9, 66.3, 64.4, 63.6, 38.6, 36.8, 35.1, 15.0; HRMS (FAB) calcd for $C_{23}H_{25}Cl_2N_3O_6S + H_1$ 542.0919, found 542.0921; Anal. Calcd for $C_{23}H_{25}Cl_2N_3O_6S$: C, 50.93; H, 4.64; N, 7.75. Found: C, 50.79; H, 4.79; N, 7.52.

Example 93.

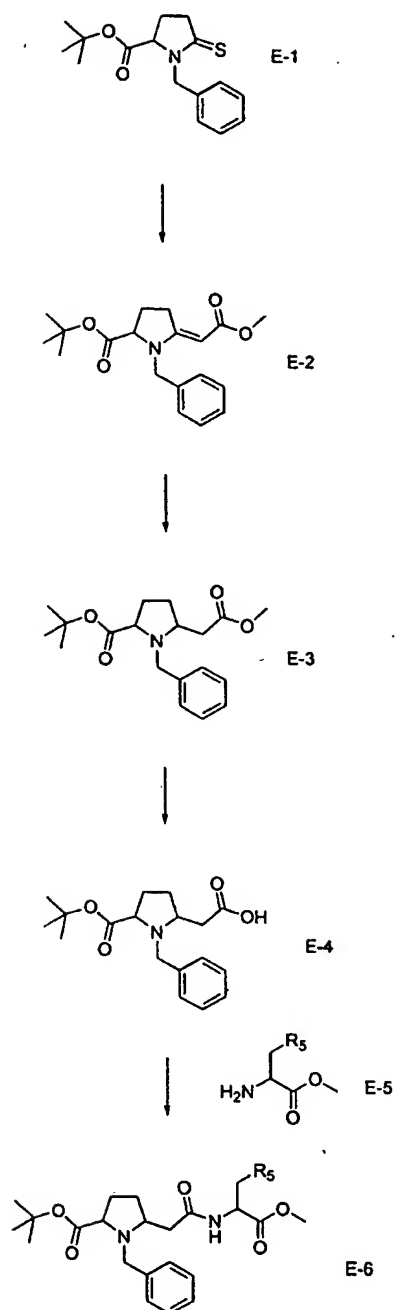
- (Scheme D, D-9: where R_{D-1} , R_{D-2} , R_{D-3} and R_{D-4} are the same and equal to proton, R_3 is 2-(1-piperidinyl)ethyl, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, Y is CO_2 and stereochemistry is (S, S)).



Example 93 was prepared as described in Scheme D from Fmoc-Tyr(2,6-Cl₂-Bn) using 1-(2-hydroxyethyl)piperidine to provide the requisite carbamate. Physical properties as follows: IR (mull) 3276, 1707, 1653, 1611, 1584, 1564, 1511, 1439, 1237, 1195, 1179, 1144, 1113, 1093, 1021 cm⁻¹; ¹H NMR (300 MHz, DMSO-d₆) δ 7.54 (2 H), 7.44 (1 H), 7.12 (2 H), 6.93 (2 H), 5.16 (2 H), 4.58 (2 H), 4.36 (1 H), 4.25 (1 H), 4.00 (3 H), 2.76 (3 H), 2.31 (3 H), 1.62 (1 H), 1.42 (3 H), 1.26 (8 H); ¹³C NMR (75 MHz, DMSO-d₆) δ 172.2, 168.9, 167.3, 157.0, 135.9, 131.7, 131.4, 130.2, 129.9, 128.7, 114.1, 64.7, 63.0, 54.0, 51.6, 36.2, 30.9, 25.5, 24.7, 23.8, 22.0, 21.5, 13.9; MS (ESI+) for C₂₈H₃₄Cl₂N₄O₆S *m/z* 624.9 (M+H)⁺.

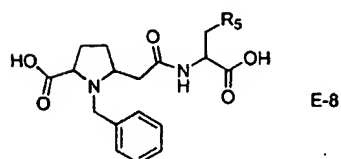
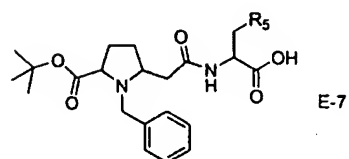
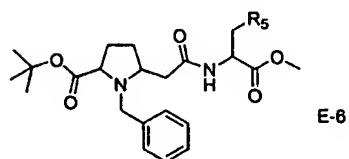
-120-

Scheme E



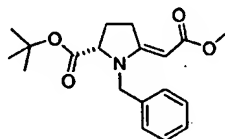
-121-

Scheme E (continued)



5

Preparation 24.

(Scheme E, E-2 where stereochemistry is (*S*)).**5- (2-Methoxy-2-oxoethylidene)-1- (phenylmethyl)-L-Proline 1,1-dimethylethyl ester**

10

To a stirring solution of E-1 (Scheme E where stereochemistry is (*S*)) (3.62g, 12.4mmol), prepared by the method of Rapoport (*J. Am. Chem. Soc.* 1984, 106, 4539), in

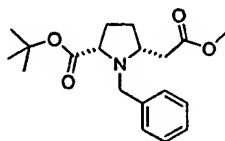
-122-

CH₃CN (10mL) was added methyl bromoacetate (1.4mL, 14.9mmol). After stirring for 70h, CH₂Cl₂ (70mL) was added. The solution stirred for 10 minutes before Ph₃P (4.89g, 18.6mmol) was added, and after 2 minutes Et₃N (5.2mL, 37.3mmol) was added. After stirring for 20h, the solution was washed with 1M NaH₂PO₄ (100mL), and the aqueous phase was extracted with CH₂Cl₂ (50mL). The combined organic phases were washed with brine, dried (Na₂SO₄), filtered, and evaporated *in vacuo*. The resulting yellow oil/white solid was dissolved in CHCl₃ and chromatographed on silica gel (300g, 230-400 mesh, 70mm OD column, packed CHCl₃, eluted with CHCl₃, 3L, then 10:90 EtOAc-CHCl₃, 250mL fractions) using the flash technique. Fractions 19-23 provided the title compound (3.23g) as a pale yellow oil. ¹H-NMR: (300MHz, CDCl₃): δ = 7.17-7.36 (5 H), 4.75 (1 H), 4.54 (1 H), 4.20 (1 H), 3.96 (1 H), 3.61 (3 H), 3.36-3.47 (1 H), 3.08 (1 H), 2.04-2.28 (2 H), 1.41 (9 H); EI/MS (70eV) *m/z* (rel. intensity): 331 (M⁺, 17.3), 275 (11.9), 230 (95.0), 170 (26.0), 91 (base); IR (neat): 2979, 2948, 1735, 1692, 1600, 1454, 1435, 1414, 1369, 1299, 1277, 1184, 1137, 1059, 964, 843, and 789cm⁻¹; HRMS: Calcd. for C₁₉H₂₅N₁O₄: 331.1783. Found: 331.1771; [α]_D²⁵: +107° (c = 0.939, CH₂Cl₂).

Preparation 25.

(Scheme E, E-3 where stereochemistry is (2R,5S)).

(2R-cis)-5-[(1,1-dimethylethoxy)carbonyl]-1- (phenylmethyl)-2-pyrrolidineacetic acid methyl ester ((2R,5S))



Raney-Nickel (20g of a 50% slurry in H₂O) was washed with abs. EtOH (3x25mL) and suspended in abs. EtOH (50mL), and a solution of E-2 (Scheme E, where

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stereochemistry is (*S*) (9.34g, 28.2mmol) in abs. EtOH (50mL) was added. After stirring for 3h, the Ra-Ni was removed by filtration, and the filtrate was evaporated *in vacuo*. The residue was dissolved in EtOAc (100mL), 5% Pt/C (3.0g) was added, and the mixture was hydrogenated under 50 psi H₂ for 12 h. The catalyst was removed by

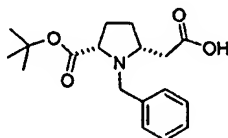
5 filtration and the filtrate was evaporated *in vacuo*. The residue was chromatographed on silica gel (300g, 230-400 mesh, 70mm OD column, packed and eluted with 15:85 EtOAc-hexanes, 270mL fractions) using the flash technique. Fractions 5-8 provided the title compound (6.55g) as a clear, colorless oil. ¹H-NMR: (300MHz, CDCl₃): δ = 7.22-7.34 (5 H), 3.86 (1 H), 3.79 (1 H), 3.62 (3 H), 3.21-3.29 (2 H), 2.57 (1 H), 2.33 (1 H),

10 1.64-2.05 (4 H), 1.37 (9 H); EI/MS (70eV) *m/z* (rel. intensity): 232 (base), 91 (39.8); IR (nujol): 2977, 1739, 1454, 1437, 1367, 1295, 1251, 1197, 1153, 1074, 844, 753, and 699cm⁻¹; Anal: Calcd. for C₁₉H₂₇N₁O₄: C, 68.44; H, 8.16; N, 4.20. Found: C, 68.39; H, 8.15; N, 4.11; [α]_D²⁵: -22° (c = 1.051, CH₂Cl₂).

Preparation 26.

15 (Scheme E, E-4 where stereochemistry is (2*R*,5*S*)).

(2*R*-cis)-5-[(1,1-dimethylethoxy)carbonyl]-1-(phenylmethyl)-2-pyrrolidineacetic acid ((2*R*,5*S*))



20

To a stirring solution of E-3 (Scheme E, where stereochemistry is (2*R*,5*S*)) (2.00g, 6.00mmol) in MeOH (60mL) was added 1*M* K₂CO₃ (20mL). After stirring for 12h, the reaction mixture was evaporated *in vacuo*, the residue was dissolved in H₂O (0.1L), the pH was adjusted to *ca.* 6 with 1*M* HCl, and the mixture was extracted with CHCl₃

25 (2X0.1L). The combined extracts were washed with H₂O, brine, dried (Na₂SO₄), and evaporated *in vacuo* to afford the title compound (1.89g) as a white solid. MP: 95-96°C

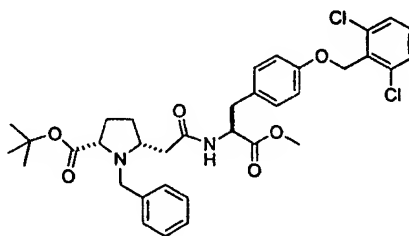
-124-

(lit. 98-101°C); ¹H-NMR: (300MHz, CDCl₃): δ = 7.30-7.37 (5 H), 4.00 (1 H), 3.68 (1 H), 3.45 (1 H), 3.20 (1 H), 2.59 (1 H), 2.46 (1 H), 1.76-2.21 (4 H), 1.31 (9 H); EI/MS (70eV) *m/z* (rel. intensity): 218 (base), 91 (86.7); IR (nujol): 1719, 1497, 1451, 1367, 1296, 1285, 1260, 1160, 1153, 1079, 965, and 757cm⁻¹; Anal: Calcd. for C₁₈H₂₅N₁O₄: C, 67.69; H, 7.89; N, 4.39. Found: C, 67.55; H, 7.97; N, 4.15; [α]_D²⁵: +27° (c = 0.795, CH₂Cl₂).

Preparation 27.

(Scheme E, E-6 where R₃ is 4-[(2,6-dichlorophenyl)methoxy]phenyl, and stereochemistry of the pyrrolidine ring is (2R,5S) and the amino acid is (S)).

(5R)-5-[2-[[(1S)-1-Methoxycarbonyl-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]-2-oxoethyl]-1- (phenylmethyl)-L-proline 1,1-dimethylethyl ester ((1S, 5R, L))



15

To a stirring solution of E-4 (Scheme E, where stereochemistry is (2R,5S)) (0.48g, 1.50mmol) in CH₂Cl₂ (10mL) was added 1- (3-dimethylaminopropyl)-3-ethylcarbodiimide hydrochloride (0.29g, 1.50mmol), 1-hydroxybenzotriazole hydrate (0.20g, 1.50mmol), 4-dimethylaminopyridine (0.05g, 0.45mmol), and 2,6-dichlorobenzyl-L-tyrosine methyl ester hydrochloride (Scheme E, E-5: where R₃ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S)) (0.59g, 1.50mmol) to give a heterogeneous mixture. Upon addition of triethylamine (0.3mL) the reaction

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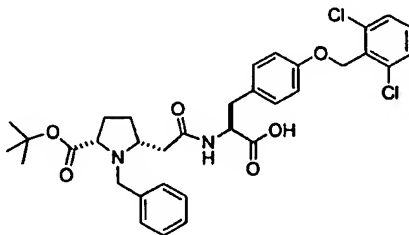
mixture became homogeneous and stirred for 12h. The reaction mixture was partitioned
 between CH₂Cl₂ (50mL) and 1N HCl (50mL). The organic phase was washed with sat'd
 aq. NaHCO₃, H₂O, brine, dried (Na₂SO₄), filtered and evaporated *in vacuo*. The
 resulting yellow oil was chromatographed on silica gel (150g, 230-400 mesh, 70mm OD
 5 column, packed and eluted acetone/CH₂Cl₂ 5:95, 40mL fractions). Fractions 36-48
 furnished the title compound (0.90g) as a glass. ¹H-NMR: (300MHz, CDCl₃): δ = 9.09 (1
 H), 7.38 (2 H), 7.10-7.30 (8 H), 6.97 (2 H), 5.22 (2 H), 4.75 (1 H), 3.81 (1 H), 3.68 (3 H),
 3.60 (1 H), 3.15-3.35 (3 H), 3.04 (1 H), 2.42 (1 H), 2.21 (1 H), 1.86-2.07 (4 H), 1.37 (9 H);
 FAB/MS *m/z* (rel. intensity): 655 (M+H, 46.4), 599 (11.3), 553 (23.6), 260 (18.8), 204
 10 (91.4), 91 (base); IR (nujol): 3262, 3001, 1733, 1665, 1612, 1585, 1565, 1512, 1439, 1392,
 1240, 1226, 1197, 1177, 1153, 1018, and 768 cm⁻¹; Anal: Calcd. for C₃₅H₄₀N₂O₆Cl₂:
 C, 64.12; H, 6.15; N, 4.27; Cl, 10.82. Found: C, 63.75; H, 6.29; N, 4.11; Cl, 10.88;
 [α]_D²⁵: +6° (c = 0.863, CHCl₃).

Preparation 28.

15 (Scheme E, E-7 where R₁ is 4-[(2,6-dichlorophenyl)methoxy]phenyl, and stereochemistry
 of the pyrrolidine ring is (2R,5S) and the amino acid is (S)).

(5R)-5-[2-[(1S)-1-Carboxy-2-[4-[(2,6-
 dichlorophenyl)methoxy]phenyl]ethyl]amino]-2-oxoethyl]-1-(phenylmethyl)-L-
 proline 1,1-dimethylethyl ester ((1S, 5R, L))

20



To a stirring solution of E-6 (Scheme E, where R₁ is 4-[(2,6-dichlorophenyl)methoxy]-
 phenyl, and stereochemistry of the pyrrolidine ring is (2R,5S) and the amino acid is (S))

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(1.00g, 1.53mmol) in MeOH (50mL) was added 1M K₂CO₃ (10mL). After stirring for 12h, the reaction mixture was evaporated *in vacuo*, the residue was dissolved in H₂O (0.1L), the pH was adjusted to *ca.* 6 with 1M HCl, and the mixture was extracted with CHCl₃ (2X0.1L). The combined extracts were washed with H₂O, brine, dried

5 (Na₂SO₄), and evaporated *in vacuo* to afford the title compound (0.85g) as a white solid.

MP: 80-83°C; ¹H-NMR: (300MHz, CDCl₃): δ = 9.87 (1 H), 7.35 (2 H), 7.21-7.26 (7 H), 7.13 (2 H), 6.94 (2 H), 5.22 (2 H), 4.64 (1 H), 3.80 (1 H), 3.55 (1 H), 3.29-3.37 (3 H), 3.04 (1 H), 2.48 (1 H), 2.26 (1 H), 1.94 (1 H), 1.76 (1 H), 1.59 (1 H), 1.41 (1 H), 1.34 (9 H);

FAB/MS *m/z* (rel. intensity): 641 (M+H, 45.8), 585 (13.0), 260 (6.1), 204 (base), 91

10 (96.8); IR (nujol): 1732, 1642, 1612, 1585, 1565, 1534, 1511, 1439, 1240, 1230, 1196, 1178, 1153, 1018, 779, and 767cm⁻¹; HRMS: Calcd. for C₃₄H₃₈Cl₂N₂O₆: 641.2185. Found. 641.2164; [α]_D²⁵: +5° (c = 0.795, CHCl₃).

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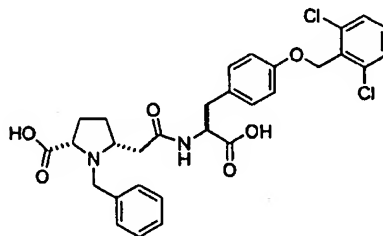
Preparation 29 and Example 94.

(Scheme E, E-8 where R₃ is 4-[(2,6-dichlorophenyl)methoxy]phenyl, and stereochemistry of the pyrrolidine ring is (2R,5S) and the amino acid is (S)).

(5R)-5-[2-[(1S)-1-Carboxy-2-[4-[(2,6-

dichlorophenyl)methoxy]phenyl]ethyl]amino]-2-oxoethyl]-1-(phenylmethyl)-L-

20 proline ((1S, 5R, L))



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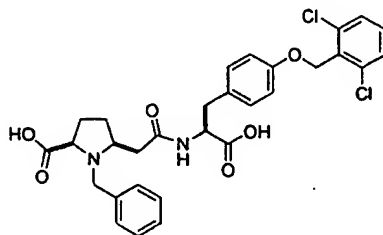
To a solution of E-7 (Scheme E, where R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl, and stereochemistry of the pyrrolidine ring is (2R,5S) and the amino acid is (S)) (0.85g, 1.32mmol) in H₂O/n-PrOH (1:1, 0.1L) was added HOAc (6mL), and the solution refluxed for 5h, then stirred at RT for 12h. Evaporation *in vacuo* afforded the title

5 compound (0.78g) as a white solid. MP: 198-202°C; ¹H-NMR: (300MHz, DMSO): δ = 8.56 (1 H), 7.56-7.59 (2 H), 7.45-7.50 (1 H), 7.25-7.29 (5 H), 7.21 (2 H), 6.96 (2 H), 5.18 (2 H), 4.41 (1 H), 3.89 (1 H), 3.74 (1 H), 3.31 (1 H), 3.03-3.09 (2 H), 2.85 (1 H), 2.18-2.34 (2 H), 1.88-2.00 (1 H), 1.63-1.84 (2 H), 1.44-1.58 (1 H); FAB/MS *m/z* (rel. intensity): 585 (M+H, 21.6), 539 (2.1), 246 (14.5), 204 (60.9), 159 (12.6), 91 (base); IR (nujol): 3309, 10
3083, 3037, 3014, 1662, 1644, 1562, 1514, 1440, 1377, 1348, 1241, 1197, 1178, 1018, 998, 815, and 771cm⁻¹; Anal: Calcd. for C₃₀H₃₀N₂O₆Cl₂O.38H₂O: C, 60.83; H, 5.24; N, 4.73. Found: C, 60.83; H, 5.33; N, 4.69; Karl Fischer water: 0.42%.

Example 95.

(Scheme E, E-8 where R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl, and stereochemistry
15 of the pyrrolidine ring is (2S,5R) and the amino acid is (S)).

(5S)-5-[2-[[(1S)-1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]-2-oxoethyl]-1- (phenylmethyl)-D-proline ((1S, 5S, D))



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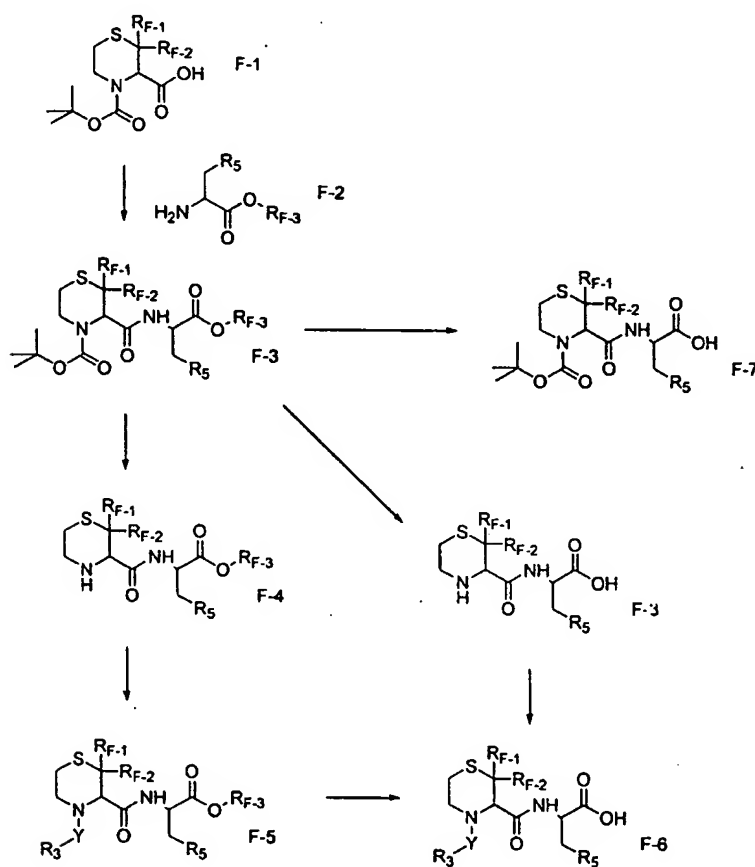
Example 95 was prepared as described in Scheme E from E-1 (Scheme E where stereochemistry is (R) prepared by the method of Rapoport (*J. Am. Chem. Soc.* 1984, 106, 4539). Physical data as follows: 198-204°C; ¹H-NMR: (300MHz, DMSO): δ = 8.60 (1 H), 7.54-7.68 (3 H), 7.26-7.41 (8 H), 7.05 (2 H), 5.27 (2 H), 4.57 (1 H), 3.96 (1 H), 3.82 (1

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H), 3.40 (1 H), 3.14 (2 H), 2.86-2.94 (1 H), 2.27-2.47 (2 H), 1.96-2.10 (1 H), 1.72-1.88 (1 H), 1.55-1.72 (1 H), 1.30-1.42 (1 H); FAB/MS m/z (rel. intensity): 585 (M+H, 50.5), 539 (4.5), 332 (32.1), 331 (18.1), 246 (6.2), 244 (6.0), 204 (77.2), 91 (base); IR (nujol): 3211, 3033, 3006, 1724, 1647, 1610, 1565, 1512, 1438, 1354, 1301, 1273, 1240, 1196, 1018, 871, and 767 cm^{-1} ; Anal: Calcd. for $\text{C}_{30}\text{H}_{30}\text{N}_2\text{O}_6\text{Cl}_2 \cdot 0.43\text{H}_2\text{O}$: C, 60.77; H, 5.24; N, 4.72.

5 Found: C, 60.76; H, 5.37; N, 4.59; Karl Fischer water analysis: 1.71%.

Scheme F



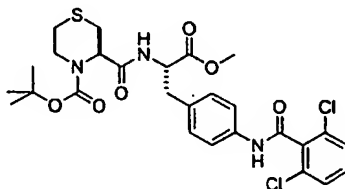
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Where $\text{R}_{\text{F-3}}$ is defined as proton or C_{1-6} alkyl.

Scheme F describes a general method for the preparation of examples of the formula F-4, F-5, F-6, F-7, and F-8. A commercially available or readily prepared sulfur containing amino acid of structure F-1 is condensed with amino acid derivative F-2 under standard peptide synthesis conditions as described in Scheme A. Deprotection of the carbamate from F-3 provides the useful intermediate F-4. The amine group may be reacted with a variety of electrophilic reagents as described in Scheme A to provide esters of general structure F-5. Mild base hydrolysis provides acids of structure F-6. Mild hydrolysis of esters of general structure F-3 provides acid of formula F-7. In those cases in which R_{F,3} is equal to *t*-butyl, mild acidolysis of compounds of general structure F-3 afford the amino acid of general structure F-8.

Preparation 30 and Example 96.

(Scheme F, F-3: where R_{F,1} and R_{F,2} are the same and equal to proton, R_{F,3} is CH₃, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*R*, *S*))



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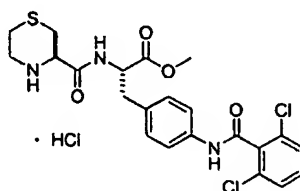
To a cooled (0 °C) solution of Boc-L-thiomorpholine-3-carboxylic acid ((a) Van Der Auwera, C.; Anteunis, M.J.O. *Int. J. Peptide Protein Res.* **1987**, *29*, 574; (b) Kogami, Y.; Okawa, K. *Bull. Chem. Soc. Jpn.* **1987**, *60*, 2963; (c) Larsson U.; Carlson R. *ACTA Chemica. Scand.* **1994**, *48*, 517; (d) Carson J.F.; Wong F.F. *J. Org. Chem.* **1964**, *29*, 2203.) (Scheme F, F-1: where R_{F,1} and R_{F,2} are the same and equal to proton and stereochemistry is (*R*)) (6.7 g, 27 mmol) in CH₂Cl₂ (100 mL) was added HOBt (4.0 g, 29.7 mmol), DMAP (700 mg), EDC (5.7 g, 29.7 mmol) and triethylamine (13.5 mL, 97 mmol). The reaction mixture was stirred for 10 min, then the amino acid derivative F-2 (Scheme F, where R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl, R_{F,3} is CH₃, and stereochemistry is (*S*)) (10.0 g, 24.7 mmol) was added. After 20 h, volatiles were removed *in vacuo* and the residue partitioned between 2.5% aqueous HCl (100 mL) and H₂O (100 mL). The organic

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layer was separated and washed saturated aqueous NaHCO₃ (100 mL), dried and concentrated *in vacuo*. Purification of the residue by chromatography on SiO₂ (500 g) using CH₂Cl₂/ethyl acetate (10%) as eluent afforded the title compound (12.31 g) as a solid: ¹H NMR (CDCl₃) δ 1.44 (9 H), 2.35 (1 H), 2.70 (3 H), 3.13 (2 H), 3.33 (1 H), 3.77 (3 H), 4.22 (1 H), 5.00 (1 H), 6.48 (1 H), 7.18 (2 H), 7.31 (3 H), 7.44 (1 H), 7.56 (2 H); ¹³C NMR (CDCl₃) δ 171.6, 168.9, 162.5, 136.5, 135.9, 132.4, 131.0, 130.2, 128.2, 120.5, 81.7, 77.3, 53.3, 52.6, 37.0, 28.2, 26.5; IR (mull) 3296, 2924, 1744, 1685, 1668, 1605, 1536, 1515, 1432, 1412, 1321, 1294, 1260, 1244, 1213, 1195, 1161, 798 cm⁻¹; MS (FAB) *m/z* (rel. intensity) 598 (M+H, 3), 596 (M+H, 5). Anal. Calcd for C₂₇H₃₁Cl₂N₃O₆S : C, 54.36; H, 5.24; N, 7.04. Found: C, 54.23; H, 5.24; N, 6.86. Corrected for 0.60% H₂O, found by Karl Fischer analysis.

Preparation 31 and Example 97.

(Scheme F, F-4: where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is CH₃, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*R*, *S*))

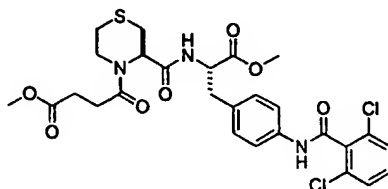


Acetyl chloride (1.75 mL, 24 mmol) was slowly added to MeOH (26 mL) at 0-5 °C. After 15 min, a solution of the carbamate F-3 (Scheme F, where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is CH₃, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*R*, *S*)) (2.4 g, 4.0 mmol) in methanol (8 mL) was added. After 50.5 h at 0 °C, the solvent was removed *in vacuo* to yield the title compound (2.11 g): ¹H NMR (300 MHz, DMSO-d₆) δ 2.68 (1 H), 3.00 (6 H), 3.47 (1 H), 3.64 (3 H), 4.02 (1 H), 4.52 (1 H), 7.28 (2 H), 7.54 (5 H), 9.15 (1 H), 9.3 (1 H), 9.70 (1 H), 10.7 (1 H); IR (mull) 3191, 3031, 1742, 1664, 1604, 1577, 1561, 1540, 1516, 1432, 1414, 1326, 1271, 1210, 799 cm⁻¹; MS (EI) *m/z* (rel. intensity) 495 (M⁺, 1).

Preparation 32 and Example 98.

(Scheme F, F-5: where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is CH_3 , R_3 is $CH_2CH_2CO_2CH_3$, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is $CO-$, and stereochemistry is (*R*, *S*)).

5



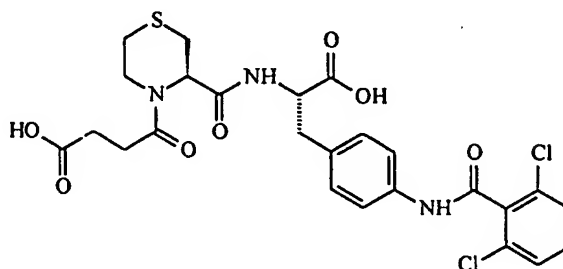
To a solution of amine F-4 (Scheme F, where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is CH_3 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*R*, *S*)) (650 mg, 1.2 mmol) was added *mono*-methyl succinate (320 mg, 2.4 mmol), EDC (460 mg, 2.4 mmol), pyridine (10 mL) and DMAP at ambient temperature. After 27 h, the mixture was diluted with 25 mL of saturated $NaHCO_3$ was extracted with methylene chloride. The combined organic extracts were dried and concentrated *in vacuo*. Purification of the residue by flash chromatography using methylene chloride/ethyl acetate (3:2) as eluant followed by lyophilization afforded the title compound (600 mg) as an amorphous solid: IR (mull) 1742, 1659, 1657, 1608, 1537, 1516, 1432, 1414, 1324, 1269, 1259, 1228, 1214, 1195, 1176 cm^{-1} ; 1H NMR (300 MHz, $CDCl_3$) δ 2.63 (7 H), 3.21 (3 H), 3.68 (3 H), 3.78 (3 H), 3.90 (1 H), 4.80 (2 H), 5.50 (1 H), 6.56 (1 H), 7.29 (5 H), 7.57 (2 H); ^{13}C NMR (75 MHz, $CDCl_3$) δ 173.9, 172.0, 171.7, 168.5, 162.7, 136.1, 135.9, 133.3, 132.3, 131.0, 130.1, 128.2, 121.0, 120.7, 53.1, 52.5, 52.3, 52.0, 44.4, 36.7, 29.4, 27.7, 26.9, 26.4; MS (EI) m/z (rel. intensity) 609 (M^+ , 5). Anal. Calcd for $C_{27}H_{29}Cl_2N_3O_7S$: C, 53.12; H, 4.79; N, 6.88. Found: C, 53.04; H, 4.81; N, 6.83. Corrected for 0.74% H_2O found by Karl Fischer analysis.

25

Preparation 33 and Example 99.

[*R*-(*R**,*S**)]- 3-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]- γ -oxo-4-thiomorpholinebutanoic acid

(Scheme F, F-6: where $R_{F,1}$ and $R_{F,2}$ are the same and equal to proton, $R_{F,3}$ is proton, R_3 is $\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is $\text{CO}-$, and stereochemistry is (R, S)).



5

To a solution of the diester F-5 (Scheme F, where $R_{F,1}$ and $R_{F,2}$ are the same and equal to proton, $R_{F,3}$ is CH_3 , R_3 is $\text{CH}_2\text{CH}_2\text{CO}_2\text{CH}_3$, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is $\text{CO}-$, and stereochemistry is (R, S)) (490 mg, 0.80 mmol) in THF (20 mL) and MeOH (6 mL) was added a solution of $\text{LiOH}\cdot\text{H}_2\text{O}$ (178 mg, 4.25 mmol) in H_2O (6 mL). After 22 h, the mixture was concentrated *in vacuo*. The residue was partially dissolved in 10% HCl (20 mL) and the resulting solid collected by filtration. The solid was washed with water and lyophilized from aqueous acetonitrile to afford the title compound (400 mg) as an amorphous solid: IR (mull) 3267, 3193, 3058, 3034, 2924, 1725, 1658, 1607, 1562, 1537, 1516, 1432, 1414, 1326, 1195, 1178, 800 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 2.62 (7 H), 3.17 (3 H), 3.95 (1 H), 4.57 (2 H), 5.26 (1 H), 7.18 (2 H), 7.55 (5 H), 8.02 (1 H), 10.64 (1 H), 12.34 (1 H); MS (FAB) m/z (rel. intensity) 582 (M+H, 18).

15

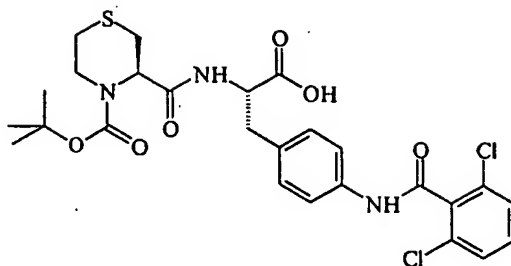
Preparation 34 and Example 100.

[R-(R*,S*)]-3-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-4-thiomorpholinecarboxylic acid 4-(1,1-dimethylethyl) ester

20

(Scheme F, F-7: where $R_{F,1}$ and $R_{F,2}$ are the same and equal to proton, $R_{F,3}$ is proton, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (R, S)).

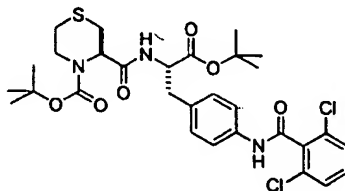
-133-



To a solution of ester **F-3** (Scheme F, where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is CH_3 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*R*, *S*)) (656 mg, 1.02 mmol) in MeOH (25 mL) was added K_2CO_3 (550 mg, 4 mmol) and H_2O (13 mL). After 3 h, volatiles were partially removed and the solution diluted with 10% HCl (20 mL) causing precipitation of a solid. The product was collected by filtration, washed with H_2O and dried in a vacuum oven to afford the product (610 mg): IR (mull) 1736, 1665, 1606, 1562, 1537, 1516, 1432, 1413, 1323, 1294, 1260, 1244, 1211, 1195, 1160 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 1.35 (9 H), 2.45 (2 H), 2.74 (5 H), 4.00 (1 H), 4.47 (1 H), 4.70 (1 H), 7.19 (2 H), 7.52 (5 H), 7.92 (1 H), 10.60 (1 H), 12.75 (1 H); MS (FAB) m/z (rel. intensity) 582 ($\text{M}+\text{H}$, 12). Anal. Calcd for $\text{C}_{26}\text{H}_{29}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$: C, 53.61; H, 5.02; N, 7.21;. Found: C, 53.20; H, 5.12; N, 7.10. Corrected for 2.30% H_2O found by Karl Fischer analysis.

Preparation 35.

(Scheme F, **F-3**: where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is *t*-butyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*R*, *S*)).



To a cooled (0-5 °C) solution of Boc-L-thiomorpholine-3-carboxylic acid (Scheme F, **F-1**: where R_{F-1} and R_{F-2} are the same and equal to proton and stereochemistry is (*R*)) (1.34 g, 5.4 mmol) in methylene chloride (20 mL) was added HOBt (800 mg, 5.94 mmol), DMAP

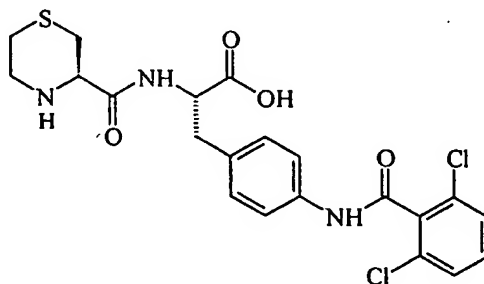
(140 mg, EDC (1.14 g, 5.94 mmol) and triethylamine (2.7 mL, 19.4 mmol). After 10 min, F-2 (Scheme F where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is *t*-butyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*)) (2.02 g, 4.94 mmol) was added, the reaction allowed to warm to ambient temperature and stirred for 24 h.

- 5 Volatiles were removed in vacuo and the residue partitioned between methylene chloride and 2.5% aqueous HCl. The organic layer was separated and washed with sat. aqueous NaHCO_3 , dried and concentrated in vacuo. Purification of the residue by flash chromatography using methylene chloride/ethyl acetate (5%) as eluant afforded the title compound (1.64 g): IR (mull) 1730, 1687, 1667, 1606, 1538, 1515, 1431, 1412, 1395, 10
1320, 1294, 1258, 1250, 1194, 1158 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 1.44 (9 H), 1.46 (9 H), 2.35 (1 H), 2.67 (3 H), 3.22 (3 H), 4.25 (1 H), 4.73 (1 H), 4.97 (1 H), 6.52 (1 H), 7.29 (5 H), 7.53 (3 H); MS (FAB) m/z (rel. intensity) 638 ($\text{M}+\text{H}$, 2). Anal. Calcd for $\text{C}_{30}\text{H}_{37}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$: C, 56.42; H, 5.84; N, 6.58. Found: C, 56.13; H, 5.98; N, 6.58.

Preparation 36 and Example 101.

- 15 4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(3R)-3-thiomorpholinyl]carbonyl]-L-phenylalanine monohydrochloride

(Scheme F, F-8: where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is proton, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*R*, *S*)).



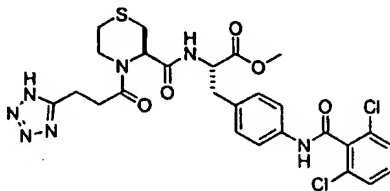
- 20 To a solution of HCl saturated in ethyl ether (5 mL) at ambient temperature was added carbamate F-3 (Scheme F, where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is *t*-butyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*R*, *S*)) (100 mg, 0.15 mmol) with vigorous stirring. After 43.5 h, the precipitate was collected by filtration and washed with ethyl ether to afford the title compound (90 mg): IR (mull) 3241, 3189,
25 3033, 2731, 1725, 1661, 1605, 1578, 1562, 1542, 1515, 1432, 1414, 1328, 1195 cm^{-1} ; ^1H

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NMR (300 MHz, DMSO- d_6) δ 2.66 (1 H), 2.99 (6 H), 3.50 (1 H), 3.98 (1 H), 4.45 (1 H), 7.26 (1 H), 7.52 (5 H), 9.00 (2 H), 9.30 (1 H), 10.69 (1 H); MS (FAB) m/z (rel. intensity) 482 (M+H, 83), 540 (32), 539 (9), 538 (42), 486 (13), 485 (16), 484 (60), 483 (24), 482 (83), 173 (11), 102 (99). Anal. Calcd for $C_{21}H_{21}Cl_2N_3O_4S \cdot HCl$: C, 48.61; H, 4.27; N, 8.10; Cl, 20.50; S, 6.18. Found: C, 48.92; H, 4.27; N, 7.79; Cl, 19.68. Corrected for 6.53% H_2O found by Karl Fischer analysis.

Example 102.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(3R)-4-[1-oxo-3-(1H-tetrazol-5-yl)propyl]-3-thiomorpholinyl]carbonyl]-L-phenylalanine methyl ester
(Scheme F, F-5: where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is CH_3 , R_3 is 2-(5-1H-tetrazolyl)ethyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is $CO-$, and stereochemistry is (R, S)).



15

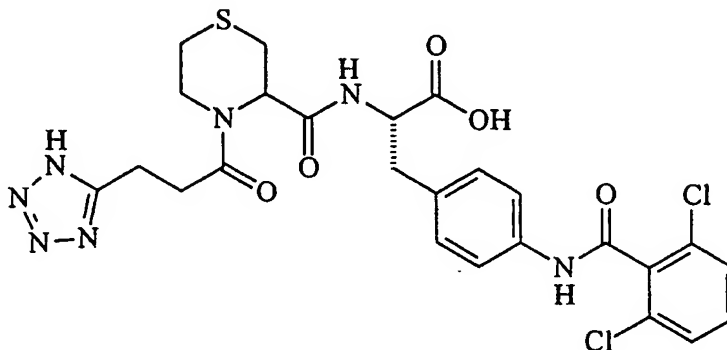
Example 102 was prepared as described in Scheme F using 1H-tetrazole-5-propanoic acid (Hutchinson, D.W.; Naylor, M. *Nucleic Acids Res.* **1985**, *13*, 8519) to form the requisite amide. Physical data as follows: IR (mull) 3264, 3047, 1742, 1659, 1607, 1561, 1537, 1516, 1432, 1415, 1324, 1268, 1219, 1195, 799 cm^{-1} ; 1H NMR (300 MHz, DMSO- d_6) δ 2.26 (1 H), 2.66 (2 H), 3.02 (8 H), 3.61 (3 H), 3.87 (1 H), 4.56 (1 H), 5.25 (1 H), 7.21 (2 H), 7.50 (5 H), 8.18 (1 H), 8.45 (1 H), 8.64 (1 H); MS (FAB) m/z (rel. intensity) 620 (M+H, 61).

25

Example 103.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(3R)-4-[1-oxo-3-(1H-tetrazol-5-yl)propyl]-3-thiomorpholinyl]-carbonyl]-L-phenylalanine

(Scheme F, F-6: where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is proton, R_3 is 2-(5-1H-tetrazolyl)ethyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is CO-, and stereochemistry is (R, S)).



5

Example 103 was prepared from example 102 by the procedure described in preparation 34. Physical data as follows: IR (mull) 3376, 3296, 3267, 3127, 1746, 1683, 1669, 1641, 1623, 1610, 1542, 1522, 1444, 1436, 1411 cm^{-1} ; MS (ESI+) for $\text{C}_{25}\text{H}_{25}\text{Cl}_2\text{N}_7\text{O}_3\text{S}$ m/z 605.8 ($\text{M}+\text{H}^+$); MS (FAB) m/z (rel. intensity) 606 (MH^+ , 51), 682 (17), 608 (40), 607 (28), 606 (51), 605 (16), 254 (99), 226 (23), 175 (17), 137 (20), 102 (33); HRMS (FAB) calcd for $\text{C}_{25}\text{H}_{25}\text{Cl}_2\text{N}_7\text{O}_3\text{S} + \text{H}_1$ 606.1093, found 606.1105.

10

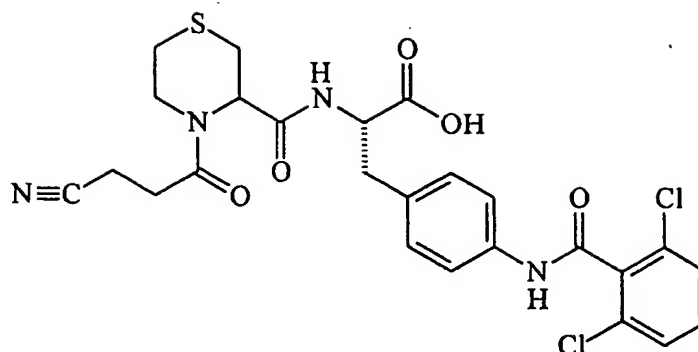
Example 104.

N-[[[(3R)-4-(3-Cyano-1-oxopropyl)-3-thiomorpholinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine

15

(Scheme F, F-6: where R_{F-1} and R_{F-2} are the same and equal to proton, R_{F-3} is proton, R_3 is $\text{CH}_2\text{CH}_2\text{CN}$, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is CO-, and stereochemistry is (R, S)).

-137-



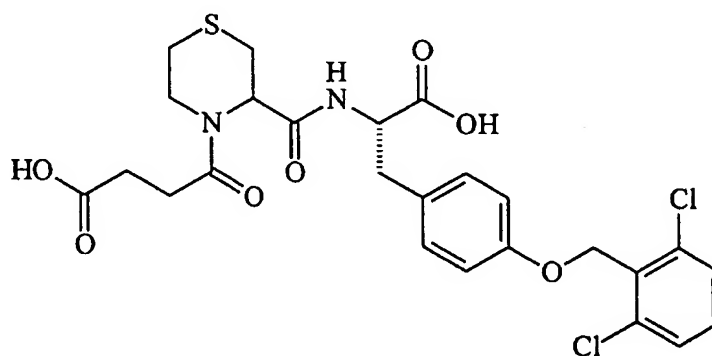
Example 104 was prepared as described in Scheme F using 3-cyanopropanoic acid (readily prepared from commercially available 3-cyanopropanoic acid) to form the requisite amide.

Physical data as follows: IR (mull) 2251, 1735, 1655, 1612, 1585, 1565, 1512, 1439,
 5 1298, 1240, 1196, 1179, 1016, 1000, 779, 768 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 2.40 (12 H), 4.88 (1 H), 5.22 (2 H), 6.72 (1 H), 6.96 (2 H), 7.23 (5 H).

Example 105.

[R-(R*,S*)]-3-[[[1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]-phenyl]ethyl]amino]carbonyl]- γ -oxo-4-thiomorpholinebutanoic acid

10 (Scheme F, F-6: where $R_{F,1}$ and $R_{F,2}$ are the same and equal to proton, $R_{F,3}$ is proton, R_3 is $\text{CH}_2\text{CH}_2\text{CO}_2\text{H}$, R_4 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, Y is CO_2 , and stereochemistry is (R, S)).

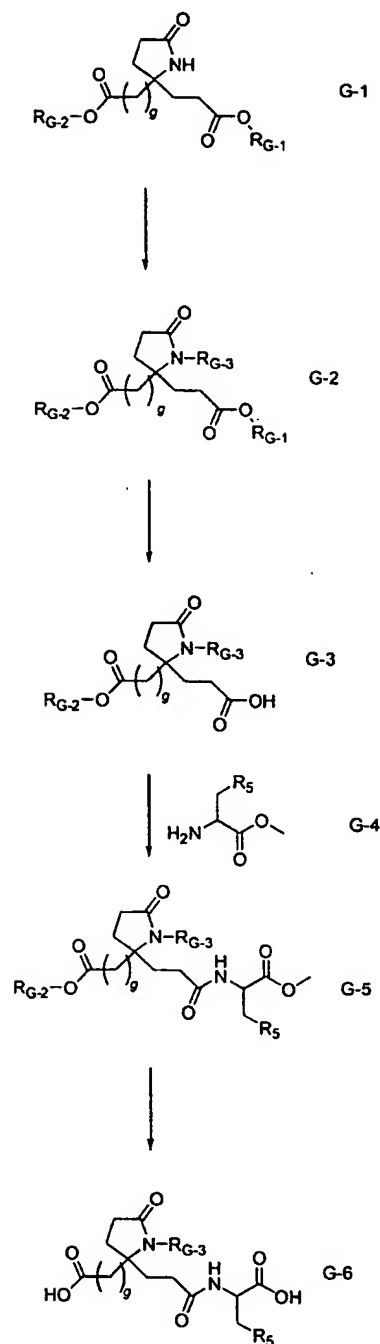


Example 105 was prepared as described in Scheme F using *mono*-methyl succinate to form
 15 the requisite amide. Physical data as follows: IR (mull) 3031, 1726, 1646, 1612, 1585, 1565, 1511, 1439, 1421, 1297, 1240, 1196, 1179, 1016, 768 cm^{-1} ; ^1H NMR (300 MHz,

DMSO- d_6) δ 2.62 (8 H), 3.64 (3 H), 4.39 (2 H), 5.20 (2 H), 6.92 (2 H), 7.15 (2 H), 7.50 (3 H), 7.98 (2 H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 174.5, 173.2, 171.9, 168.9, 136.5, 132.3, 132.0, 130.8, 130.7, 129.2, 114.8, 65.3, 54.3, 52.5, 36.0, 29.7, 28.0, 27.0; MS (FAB) m/z (rel. intensity) 569 (M+H, 24). Anal. Calcd for $\text{C}_{23}\text{H}_{26}\text{Cl}_2\text{N}_2\text{O}_7\text{S}$: C, 52.73; H, 4.60; N, 4.92; Cl, 12.45. Found: C, 52.51; H, 4.60; N, 4.94; Cl, 12.78. Corrected for 3.37% H_2O found by Karl Fischer analysis.

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Scheme G



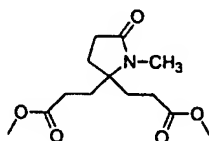
Where R_{G-1} and R_{G-2} are defined independently as H or CH_3 ; R_{G-3} is defined as H, C_{1-6} alkyl or C_{3-6} alkenyl; and g is defined as 0 or 2.

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Scheme G describes a general method to prepare lactam examples of general structures G-5 and G-6. Readily prepared lactams of general structure G-1 may be alkylated by the reaction of an appropriate alkylating in the presence of a suitable base as described in preparation 37 to provide intermediates of general structure G-2. Mild hydrolysis provides the monoacid of general structure G-3 which may be condensed with an amino acyl intermediate of structure G-4 as described in Scheme A. Full hydrolysis of the diester of general structure G-5 affords the diacid of structure G-6.

Preparation 37.

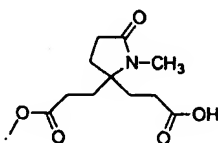
(Scheme G, G-2: where R_{G-1} and R_{G-2} are CH_3 , R_{G-3} is CH_3 and g is equal to 2).



To a cooled (0-5 °C) solution of dimethyl ester G-1 (Scheme G, where R_{G-1} and R_{G-2} are CH_3 , R_{G-3} is H and g is equal to 2) (Thomas, E.T.; Rynbrandt, R.H; Zimmermann, D.C.; Bell, L.T.; Muchmore, C.R.; Yankee, E.W. *J. Org. Chem.* 1989, 54, 4535) (25.7 g, 0.1 mole) and iodomethane (30 mL, 0.48 mol) in THF (200 mL) was added NaH (4.8 g, 0.12 mmol, 60% in oil dispersion). After 22 h, the reaction was quenched by the addition of H_2O (100 mL) and diluted with CH_2Cl_2 (50 mL). The organic layer was separated, dried and dried *in vacuo*. The crude brown oil was triturated with hexanes (200 mL) and the residue concentrated *in vacuo* to afford the crude product (18.44 g) which was used without further purification: 1H NMR (300 MHz, $CDCl_3$) δ 2.03 (12 H), 2.64 (3 H), 3.64 (6 H).

Preparation 38.

(Scheme G, G-3: where R_{G-2} is CH_3 , R_{G-3} is CH_3 and g is equal to 2).



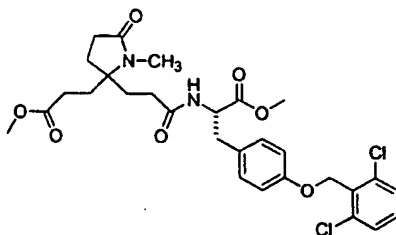
-141-

To a cooled (0-5 °C) solution of diester **G-2** (Scheme G, where R_{G-1} and R_{G-2} are CH_3 , R_{G-3} is CH_3 and g is equal to 2) (10.0 g, 36.9 mmol) in aqueous methanol (66%, 75 mL) was added $LiOH \cdot H_2O$ (1.55 g, 36.9 mmol). After 22 h, the mixture was partially concentrated
 5 *in vacuo* and diluted with water. The aqueous layer was washed with methylene chloride and acidified (pH ca. 2) with 10% HCl. The aqueous layer was extracted with methylene chloride and the combined organic extracts dried and concentrated *in vacuo* to afford the title compound (3.63 g) as a yellow solid: IR (mull) 1735, 1630, 1442, 1429, 1418, 1405, 1330, 1308, 1287, 1255, 1217, 1185, 1122, 1016, 642 cm^{-1} ; 1H NMR (300 MHz, $CDCl_3$) δ
 10 1.90 (6 H), 2.24 (4 H), 2.43 (2 H), 2.68 (3 H), 3.67 (3 H); ^{13}C NMR (75 MHz, $CDCl_3$) δ 176.3, 175.5, 173.2, 64.6, 52.0, 33.1, 32.9, 29.9, 28.41, 28.38, 26.0, 24.9; MS (EI) m/z (rel. intensity) 257 (M^+ , 1).

Preparation 39 and Example 106.

2-[3-[[[(1S)-1-[[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]-3-oxopropyl]-1-methyl-5-oxo-2-pyrrolidinepropanoic acid methyl
 15 ester

(Scheme G, **G-5**: where R_{G-2} and R_{G-3} are both equal to CH_3 , R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, g is equal to 2 and stereochemistry is (*S*)).



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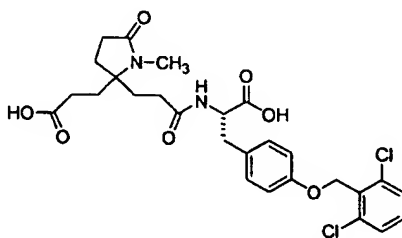
To a solution of acid **G-3** (Scheme G, where R_{G-2} and R_{G-3} are equal to CH_3 and g is equal to 2) (1.0 g, 3.9 mmol), amine **G-4** (Scheme G, where R_5 is 4-[(2,6-dichlorophenyl)-methoxy]phenyl and stereochemistry is (*S*)) (1.82 g, 4.67 mmol), and DMAP (100 mg, 0.8
 25 mmol) in pyridine (15 mL) was added EDC (895 mg, 4.67 mmol). After 21 h, the reaction was diluted with saturated aqueous $NaHCO_3$ and methylene chloride. The organic

layer was separated and washed with 10% aqueous HCl, dried and concentrated *in vacuo*. Purification of the residue by flash chromatography using methylene chloride/ethyl acetate (20%) as eluant followed by lyophilization from aqueous acetonitrile afforded the title compound (1.71 g) as an amorphous solid: IR (mull) 3284, 2924, 1738, 1666, 1665, 1564, 1539, 1511, 1439, 1398, 1299, 1240, 1198, 1178, 1117, 1016, 768 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 2.10 (12 H), 2.66 (3 H), 3.07 (2 H), 3.66 (3 H), 3.73 (3 H), 4.83 (1 H), 5.24 (2 H), 5.98 (1 H), 6.95 (2 H), 7.02 (2 H), 7.25 (1 H), 7.36 (2 H); ^{13}C NMR (75 MHz, CDCl_3) δ 174.7, 174.66, 173.2, 172.0, 171.2, 158.1, 137.0, 132.1, 130.5, 130.3, 128.5, 128.3, 115.2, 65.2, 64.3, 53.3, 52.4, 51.9, 37.0, 33.2, 30.2, 30.17, 28.4, 26.0, 24.7; MS (EI) m/z (rel. intensity) 592 (M^+ , 4). Anal. Calcd for $\text{C}_{29}\text{H}_{34}\text{Cl}_2\text{N}_2\text{O}_7$: C, 58.69; H, 5.77; N, 4.72; Cl, 11.95. Found: C, 58.33; H, 5.65; N, 4.76; Cl, 11.89.

Preparation 40 and Example 107.

2-[3-[[[(1S)-1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]-3-oxo-propyl]-1-methyl-5-oxo-2-pyrrolidinepropanoic acid

(Scheme G, G-6: where $\text{R}_{\text{G-3}}$ is equal to CH_3 , R_g is 4-[(2,6-dichlorophenyl)methoxy]phenyl, g is equal to 2 and stereochemistry is (S)).



To a solution of diester G-5 (Scheme G, where $\text{R}_{\text{G-2}}$ and $\text{R}_{\text{G-3}}$ are both equal to CH_3 , R_g is 4-[(2,6-dichlorophenyl)methoxy]phenyl, g is equal to 2 and stereochemistry is (S)) (1.0 g, 1.68 mmol), in THF (30 mL) and MeOH (9 mL) was added $\text{LiOH}\cdot\text{H}_2\text{O}$ (370 mg, 8.8 mmol). After 22.5 h, the reaction mixture was acidified with 10% aqueous HCl (30 mL) causing precipitation of a solid. The solid was collected by filtration and lyophilized from aqueous acetonitrile to afford the title compound (0.91 g) as an amorphous solid: IR (mull) 3031, 2925, 1727, 1637, 1585, 1564, 1543, 1511, 1439, 1424, 1404, 1299, 1240,

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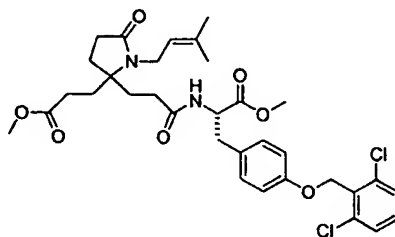
1196, 1179, 768 cm^{-1} ; ^1H NMR (300 MHz, DMSO-d_6) δ 1.77 (12 H), 2.76 (1 H), 3.97 (1 H), 4.36 (1 H), 5.16 (2 H), 6.94 (2 H), 7.15 (2 H), 7.49 (3 H), 8.16 (1 H); ^{13}C NMR (75 MHz, CDCl_3) δ 174.6, 174.1, 173.6, 172.2, 157.6, 136.5, 132.2, 132.0, 130.7, 130.6, 129.2, 114.7, 65.3, 64.1, 54.0, 36.5, 33.7, 33.1, 30.1, 30.05, 29.8, 28.7, 25.7, 24.6; MS (FAB) m/z (rel. intensity) 565 ($\text{M}+\text{H}$, 99). Anal. Calcd for $\text{C}_{27}\text{H}_{30}\text{Cl}_2\text{N}_2\text{O}_7$: C, 57.35; H, 5.35; N, 4.95; Cl, 12.54. Found: C, 56.93; H, 5.15; N, 5.02; Cl, 12.42. Corrected for 1.03% H_2O found by Karl Fischer analysis.

Example 108.

10 2-[3-[[[(1S)-1-[[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]-3-oxopropyl]-1-(3-methyl-2-butenyl)-5-oxo-2-pyrrolidinepropanoic acid methyl ester

(Scheme G, G-5: where $\text{R}_{\text{G-2}}$ is CH_3 , $\text{R}_{\text{G-3}}$ is 1-(3-methyl-2-butenyl), R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, g is equal to 2 and stereochemistry is (*S*)).

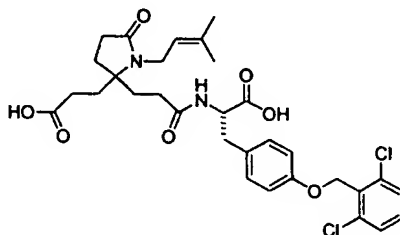
15



Example 108 was prepared as described in Scheme G using 1-bromo-3-methyl-2-butene to form the requisite N-alkyl lactam. Physical data as follows: IR (mull) 32.95, 29.52, 1740, 1678, 1662, 1564, 1538, 1512, 1439, 1414, 1300, 1279, 1240, 1199, 1178, 1017, 768 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 1.64 (3 H), 1.70 (3 H), 2.06 (12 H), 3.09 (2 H), 3.66 (3 H), 3.74 (2 H), 3.74 (3 H), 4.85 (1 H), 5.20 (1 H), 5.24 (2 H), 6.95 (2 H), 7.02 (2 H), 7.25 (1 H), 7.36 (2 H); ^{13}C NMR (75 MHz, CDCl_3) δ 174.7, 173.3, 172.1, 172.0, 171.3, 158.1, 137.0, 135.1, 132.1, 130.5, 130.3, 128.5, 128.3, 119.7, 115.2, 65.2, 65.1, 53.2, 52.4, 51.9, 37.5, 37.0, 34.4, 34.2, 30.6, 29.9, 28.6, 26.6, 25.6, 25.5, 17.9; MS (FAB) m/z (rel. intensity) 647 ($\text{M}+\text{H}$, 24).

Example 109.

2-[3-[[[(1S)-1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]-3-oxo-propyl]-1-(3-methyl-2-butenyl)-5-oxo-2-pyrrolidinepropanoic acid
(Scheme G, G-6: where R_{G-3} is 1-(3-methyl-2-butenyl), R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl, g is equal to 2 and stereochemistry is (S)).



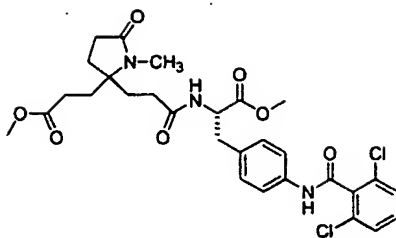
Example 109 was prepared from example 108 by the procedure described in preparation
40. Physical data as follows: IR (mull) 3290, 2921, 1726, 1635, 1585, 1565, 1545, 1511, 1439, 1419, 1341, 1299, 1240, 1197, 1179, 780, 768 cm⁻¹; ¹H NMR (300 MHz, DMSO-d₆) δ 1.57 (3 H), 1.62 (3 H), 1.72 (6 H), 2.02 (6 H), 2.77 (1 H), 2.98 (1 H), 3.55 (2 H), 5.05 (1 H), 5.16 (2 H), 6.94 (2 H), 7.15 (2 H), 7.49 (2 H), 7.54 (2 H), 8.15 (1 H); ¹³C NMR (75 MHz, DMSO-d₆) δ 174.6, 174.3, 173.6, 173.56, 172.2, 157.6, 136.5, 134.0, 132.2, 132.0, 130.6, 129.2, 121.1, 121.0, 114.7, 65.2, 65.0, 54.1, 54.0, 37.1, 36.5, 34.5, 34.1, 30.0, 29.9, 28.8, 26.3, 25.84, 25.81, 18.1; MS (FAB) m/z (rel. intensity) 619 (M+H, 99).

Example 110.

2-[3-[[[(1S)-1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]-3-oxopropyl]-1-methyl-5-oxo-2-pyrrolidinepropanoic acid methyl ester

(Scheme G, G-5: where R_{G-2} and R_{G-3} are equal to CH₃, R₅ is 4-[(2,6-dichlorobenzoyl)-amino]phenyl, g is equal to 2 and stereochemistry is (S)).

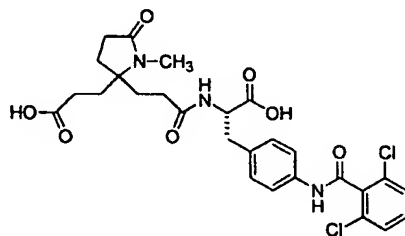
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Example 110 was prepared as described in Scheme G using iodomethane to form the requisite N-alkyl lactam. Physical data as follows: IR (mull) 3258, 2922, 1738, 1662,
 5 1606, 1561, 1539, 1515, 1432, 1414, 1401, 1323, 1268, 1196, 1177, 799 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 2.01 (12 H), 2.61 (3 H), 3.09 (2 H), 3.64 (3 H), 3.75 (3 H), 4.84 (1 H), 6.15 (1 H), 7.09 (2 H), 2.31 (3 H), 7.58 (2 H), 7.99 (1 H); MS (EI) m/z (rel. intensity) 607 (M^+ , 4), 605 (M^+ , 6); Anal. Calcd for $\text{C}_{29}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_7$: C, 57.43; H, 5.48; N, 6.93; Cl, 11.69. Found: C, 57.18; H, 5.56; N, 6.85; Cl, 11.68. Corrected for 0.93% H_2O , found by
 10 Karl Fischer analysis.

Example 111.

2-[3-[[[(1S)-1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]-3-oxo-propyl]-1-methyl-5-oxo-2-pyrrolidinepropanoic acid
 (Scheme G, G-6: where $\text{R}_{\text{G-3}}$ is CH_3 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, g is
 15 equal to 2 and stereochemistry is (S)).



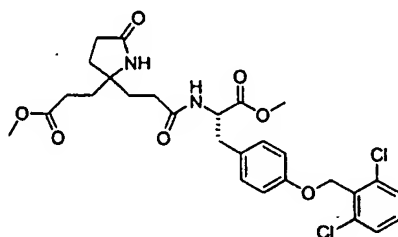
Example 111 was prepared from example 110 by the procedure described in preparation
 20 40. Physical data as follows: IR (mull) 3265, 3056, 2925, 1724, 1658, 1609, 1579, 1561, 1542, 1516, 1432, 1414, 1327, 1271, 1217, 1195, 800 cm^{-1} ; ^1H NMR (300 MHz, DMSO-

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d₆) δ 1.89 (12 H), 2.49 (3 H), 2.78 (1 H), 2.99 (1 H), 4.38 (1 H), 7.18 (2 H), 7.51 (5 H), 8.17 (1 H), 10.64 (1 H); MS (FAB) *m/z* (rel. intensity) 580.5 (M+H, 68).

Example 112.

2-[3-[[[(1S)-1-[[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]-3-oxopropyl]-5-oxo-2-pyrrolidinepropanoic acid methyl ester
 (Scheme G, G-5: where R_{G-2} is CH₃, R_{G-3} is proton, R₅ is 4-[(2,6-dichlorophenyl)-methoxy]phenyl, g is equal to 2 and stereochemistry is (S)).



10

Example 112 was prepared as described in Scheme G. Physical data as follows: IR (mull) 3276, 3029, 1738, 1686, 1564, 1538, 1511, 1439, 1299, 1279, 1239, 1197, 1178, 1016, 767, cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 1.85 (6 H), 2.24 (2 H), 2.35 (4 H), 3.03 (2 H), 3.66 (3 H), 3.74 (3 H), 4.82 (1 H), 5.24 (2 H), 6.50 (2 H), 6.95 (2 H), 7.05 (2 H), 7.27 (1 H), 7.37 (2 H); MS (EI) *m/z* (rel. intensity) 578 (M⁺, 0.2); Anal. Calcd for C₂₈H₃₂Cl₂N₂O₇: C, 58.04; H, 5.57; N, 4.83. Found: C, 57.93; H, 5.43; N, 4.97. Corrected for 1.14 % H₂O found by Karl Fischer analysis.

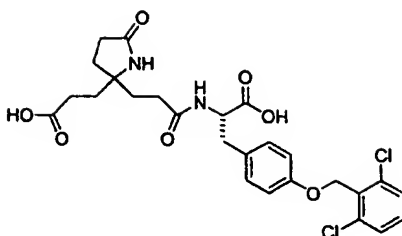
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Example 113.

2-[3-[[[(1S)-1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]-3-oxo-propyl]-5-oxo-2-pyrrolidinepropanoic acid
 (Scheme G, G-6: where R_{G-3} is proton, R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl, g is equal to 2 and stereochemistry is (S)).

20

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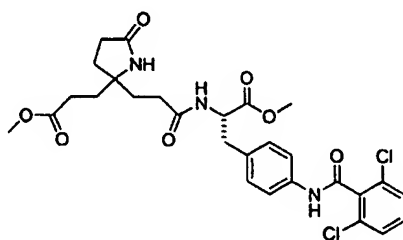


Example 113 was prepared from example 112 by the procedure described in preparation.

40. Physical data as follows: IR (mull) 3294, 3033, 1716, 1647, 1585, 1565, 1544, 1511,
 5 1439, 1420, 1299, 1240, 1197, 1179, 768 cm^{-1} ; ^1H NMR (300 MHz, DMSO-d_6) δ 1.60
 (6 H), 2.12 (6 H), 2.78 (1 H), 2.99 (1 H), 4.36 (1 H), 5.19 (2 H), 6.97 (2 H), 7.17 (2 H),
 7.45 (1 H), 7.58 (2 H), 7.70 (1 H), 8.22 (1 H); MS (FAB) m/z (rel. intensity) 551 ($\text{M}+\text{H}$,
 99); Anal. Calcd for $\text{C}_{26}\text{H}_{28}\text{Cl}_2\text{N}_2\text{O}_7$: C, 56.63; H, 5.12; Cl, 12.86; N, 5.08. Found: C,
 56.28; H, 5.01; Cl, 13.08; N, 5.24. Corrected for 1.47% H_2O found by Karl Fischer
 10 analysis.

Example 114.

2-[3-[[[(1S)-1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-
 oxoethyl]amino]-3-oxopropyl]-5-oxo-2-pyrrolidinepropanoic acid methyl ester
 (Scheme G, G-5: where $\text{R}_{\text{G}2}$ is CH_3 , $\text{R}_{\text{G}3}$ is proton, R_5 is 4-[(2,6-dichlorobenzoyl)-
 15 amino]phenyl, g is equal to 2 and stereochemistry is (*S*)).

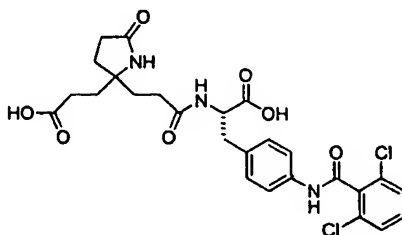


Example 114 was prepared as described in Scheme G. Physical data as follows: ^1H NMR
 20 (300 MHz, CDCl_3) δ 1.81 (6 H), 2.27 (6 H), 3.10 (2 H), 3.63 (3 H), 3.75 (3 H), 4.89 (1 H),
 6.46 (1 H), 6.58 (1 H), 7.10 (2 H), 7.26 (3 H), 7.58 (2 H), 8.14 (1 H); MS (FAB) m/z 592
 ($\text{M}+\text{H}$) $^+$, 568, 367, 349, 307, 278, 226, 194, 173.

Example 115.

2-[3-[[[(1S)-1-Carboxy-1-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]-3-oxopropyl]-5-oxo-2-pyrrolidinepropanoic acid

(Scheme G, G-6: where R_{G-3} is proton, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, g is equal to 2 and stereochemistry is (*S*)).



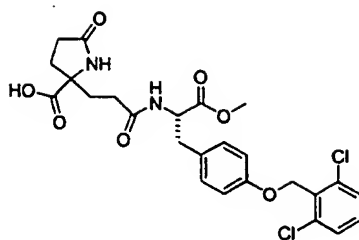
Example 115 was prepared from example 114 by the procedure described in preparation 40. Physical data as follows: IR (mull) 3272, 3195, 3121, 3063, 2953, 2923, 2868, 2855, 1715, 1659, 1608, 1579, 1561, 1541, 1516, 1456, 1432, 1414, 1377, 1367, 1326, 1271, 1221, 1195, 800 cm^{-1} ; ^1H NMR (300 MHz, DMSO- d_6) δ 1.63 (6 H), 2.13 (6 H), 2.79 (1 H), 3.00 (1 H), 4.38 (1 H), 7.19 (2 H), 7.51 (5 H), 7.64 (1 H), 8.16 (1 H), 10.58 (1 H), 12.37 (2 H); MS (FAB) m/z 564 ($M+H$) $^+$, 546, 519, 335, 280, 194, 173. Anal. Calc'd for $C_{26}H_{27}Cl_2N_3O_7$: C, 55.33; H, 4.82; Cl, 12.56; N, 7.44. Found: C, 55.10; H, 4.76; Cl, 12.56; N, 7.36. Corrected for 2.49% H_2O , found by Karl Fischer analysis.

Example 116.

2-[3-[[[(1S)-1-[[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]-3-oxopropyl]-5-oxoproline

(Scheme G, G-5: where R_{G-2} and R_{G-3} are equal to proton, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl, g is equal to 0 and stereochemistry is (*S*)).

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Example 115 was prepared as described in Scheme G from 2-carboxy-5-oxo-2-pyrrolidinepropanoic acid (Majer, Z.; Kajtar, M.; Tichy, M.; Blaha, K. *Coll. Czech. Chem. Commun.* **1982**, *47*, 950). Physical data as follows: IR (mull) 3302, 1736, 1671, 1612, 1585, 1564, 1535, 1511, 1439, 1298, 1240, 1197, 1179, 1016, 768 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 2.12 (8 H), 3.02 (2 H), 3.65 (3 H), 4.81 (1 H), 5.21 (2 H), 6.94 (2 H), 7.07 (2 H), 7.22 (1 H), 7.35 (2 H), 7.86 (1 H), 8.34 (1 H); MS (FAB) m/z (rel. intensity) 537 (M+H, 99).

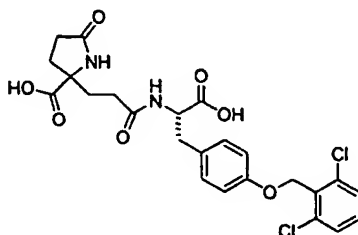
10

Example 117.

2-[3-[[[(1S)-1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]-3-oxo-propyl]-5-oxoproline

(Scheme G, G-6: where R_{G-3} is proton, R_g is 4-[(2,6-dichlorophenyl)methoxy]phenyl, g is equal to 0 and stereochemistry is (S)).

15



20

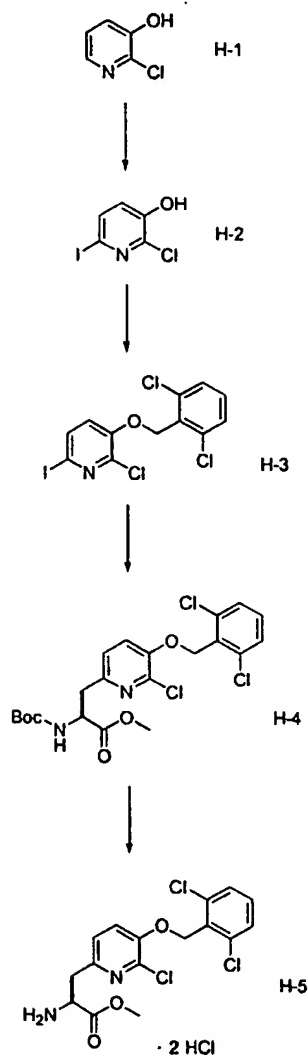
Example 116 was prepared from example 115 by the procedure described in preparation 40. Physical data as follows: IR (mull) 3292, 3059, 3029, 1718, 1650, 1612, 1585, 1565, 1537, 1511, 1439, 1240, 1196, 1179, 768 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 2.11 (8 H),

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2.83 (1 H), 3.20 (1 H), 4.74 (1 H), 5.24 (2 H), 6.95 (2 H), 7.15 (2 H), 7.25 (1 H), 7.36 (2 H); MS (FAB) m/z (rel. intensity) 523 (M+H, 99); Anal. Calcd for $C_{24}H_{24}Cl_2N_2O_7$: C, 55.08; H, 4.62; N, 5.35, Cl, 13.55. Found: C, 54.68; H, 4.66; N, 5.13; Cl, 13.70. Corrected for 1.59% H_2O found by Karl Fischer analysis.

5

Scheme H.



Scheme H teaches a general method for the preparation of 6-chloroazatyrosine examples of structures **H-4** and **H-5**, by adaption of the methodology for the preparation of protected azatyrosine reagents as described by Kawata, S.; Ashizawa, S.; Hirama, M. *J. Am. Chem. Soc.* 1997, 119, 12012-12013 and references cited therein. Thus regioselective iodination of 2-chloro-3-pyridinol gives the chloro-iodopyridinol **H-2**, that is *O*-alkylated as exemplified by the synthesis of **H-3**. Palladium-catalyzed reaction of **H-3** with the organozinc derived from a suitably protected β -iodoalanine, provides the protected 5-chloroazatyrosine **H-4**. *N*-deprotection of **H-4** gives the aminoester **H-5**, that is used (as exemplified by reagent A-4 of Scheme A, and by reagent B-5 of Scheme B) for the synthesis of Examples of this invention

Preparation 41.

(Scheme H: **H-2**)**2-Chloro-6-iodo-3-pyridinol** (C_5H_3ClINO).

To a solution of 2-chloro-3-pyridinol **H-1** (10.2 g, 78.7 mmol) and K_2CO_3 (38.9 g, 0.274 mol) in H_2O (120 mL) is added I_2 (24.3 g, 95.8 mmol), and the reaction mixture is stirred at rt for 4 h. The reaction mixture is quenched by the addition of aq satd $Na_2S_2O_3 \cdot 5H_2O$, and its pH is lowered to pH 2 with the addition of 12 M aq HCl. The mixture is extracted with EtOAc. The combined EtOAc extracts are dried, filtered and concentrated to a yellow solid, that is crystallized from 120:25 heptane/EtOAc (145 mL) to give, as a yellow solid, 11.2 g of the title compound: 1H NMR (CD_3SOCD_3 , 300 MHz) δ 9.87 (1H), 7.59 (1H), 7.06 (1H); ^{13}C NMR (CD_3SOCD_3 , 75 MHz) δ 150.68, 138.07, 134.98, 127.02, 101.18.

Preparation 42.

(Scheme H: **H-3**)**2-Chloro-3-[(2,6-dichlorophenyl)methoxy]-6-iodopyridine** ($C_{12}H_7Cl_3INO$).

To a solution of **H-2** (5.11 g, 20.0 mmol), PPh_3 (5.30 g, 20.0 mmol), and 2,6-dichlorobenzylalcohol (3.54 g, 20.0 mmol) in dry THF (100 mL) at 0 °C under Ar is added

DEAD (3.15 mL, 20.0 mmol). The reaction mixture is kept at 0 °C for 1.5 h and at rt for 1.5 h. It is concentrated to a residue, that is purified by silica flash chromatography (17:3 hexanes/EtOAc) to give 7.61 g of the title compound: TLC (17:3 hexanes/EtOAc) R_f 0.57; ^1H NMR (CD_3SOCD_3 , 300 MHz) δ 7.85 (1H), 7.62 (1H), 7.58–7.45 (3H), 5.34 (2H).

5

Preparation 43.

(Scheme H: H-4)

(*S*)-2-Chloro-3-[(2,6-dichlorophenyl)methoxy]- α -[(1,1-dimethylethoxy)-carbonyl]amino]-6-pyridinepropanoic acid methyl ester ($\text{C}_{21}\text{H}_{23}\text{Cl}_3\text{N}_2\text{O}_5$).

To an amberized flask containing activated Zn dust (0.777 g, 11.89 mmol) under Ar is added sequentially *N*-[(1,1-dimethylethoxy)carbonyl]-3-iodo-L-alanine methyl ester [93267-04-0] (3.91 g, 1.89 mmol), THF (11.9 mL), and dimethylacetamide (11.9 mL). The reaction mixture was purged of O_2 by the bubbling of Ar through the mixture for 5 min. It is stirred at 65 ± 5 °C for 2 h, and is cooled to rt. To this mixture is added $\text{PdCl}_2(\text{PPh}_3)_2$ (0.412 g), followed immediately afterward by a degassed solution of H-3 (2.46 g, 5.94 mmol) in 1:1 THF/dimethylacetamide (11.8 mL). The reaction mixture is stirred at 65 ± 5 °C for 5 h. It is cooled to 0 °C and quenched with sat'd aq NH_4Cl (100 mL). The reaction mixture is extracted with EtOAc. The combined extracts are washed with brine, dried, filtered and concentrated to a green-yellow oil; that is purified by silica flash chromatography to give 1.90 g of the title compound: TLC (7:3 hexanes/EtOAc) R_f 0.32; ^1H NMR (CDCl_3 , 300 MHz) δ 7.76 (1H), 7.57 (2H), 7.48 (1H), 7.29 (1H), 7.27 (1H), 5.30 (2H), 4.32 (1H), 3.60 (3H), 3.01 (1H), 2.98 (1H), 1.31 (9H).

15

20

Preparation 44.

(Scheme H: H-4)

(*S*)-2-Chloro-3-[(2,6-dichlorophenyl)methoxy]- α -amino-6-pyridinepropanoic acid methyl ester dihydrochloride salt ($\text{C}_{16}\text{H}_{15}\text{Cl}_3\text{N}_2\text{O}_3 \cdot 2\text{HCl}$).

25

A solution of H-4 (1.90 g, 3.88 mmol) in 4 M HCl in dioxane (35 mL) is stirred at rt under Ar for 20 h. The reaction mixture is concentrated *in vacuo*. The residue is taken up in H_2O (40 mL), and extracted with Et_2O . The aqueous solution is frozen and lyophilized

to give 1.39 g of the title compound: ^1H NMR (CD_3SOCD_3 , 300 MHz) δ 8.62 (3H), 7.81 (1H), 7.58 (2H), 7.48 (1H), 7.38 (1H), 5.32 (2H), 4.37 (1H), 3.72 (3H), 3.27 (2H).

Example 118.

(Scheme A, A-5)

- 5 [S-(*R**,*R**)]-4-[[[1-[[2-Chloro-3-[(2,6-dichlorophenyl)methoxy]-6-pyridyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid (1,1-dimethylethyl) ester ($\text{C}_{30}\text{H}_{37}\text{Cl}_3\text{N}_3\text{O}_6\text{S}$).

Example 118 was prepared as described in Scheme A from D-cysteine using the product of preparation 44 as amino acid intermediate A-4. Physical properties as follows:

- 10 TLC (1:1 hexanes/EtOAc) R_f 0.27; ^1H NMR (CDCl_3 , 300 MHz) δ 8.43 (1H), 7.74 (1H), 7.58 (2H), 7.48 (1H), 7.29 (1H), 5.29 (2H), 4.67 (1H), 4.53 (1H), 4.44 (1H), 4.23 (1H), 3.62 (3H), 3.06 (3H), 2.82 (1H), 1.27 (9H).

Example 119.

(Scheme A, A-9)

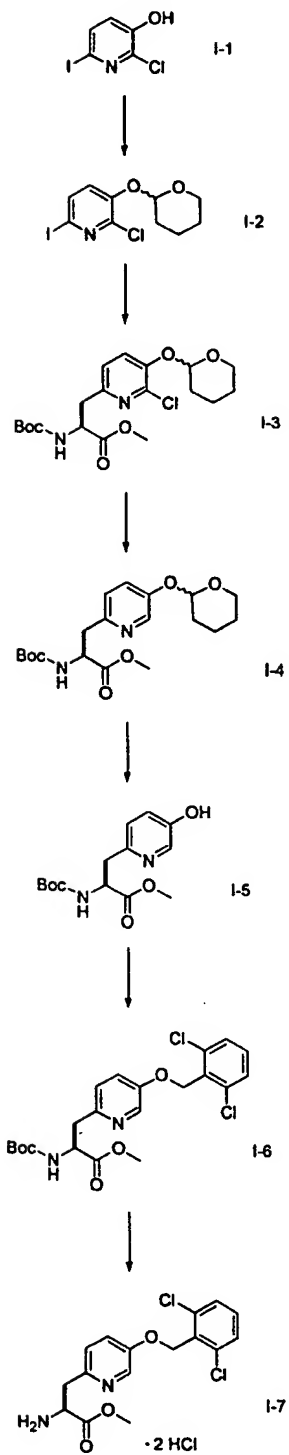
- 15 [S-(*R**,*R**)]-4-[[[1-Carboxy-2-[[2-Chloro-3-[(2,6-dichlorophenyl)methoxy]-6-pyridyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid (1,1-dimethylethyl) ester ($\text{C}_{24}\text{H}_{26}\text{Cl}_3\text{N}_3\text{O}_6\text{S}$).

Example 119 was prepared from example 118 by the procedure described in preparation

12. Physical data as follows: mp 93–95 °C; TLC (600:400:2 EtOAc/hexanes/ HCO_2H) R_f
20 0.38; ^1H NMR (CD_3SOCD_3 , 300 MHz) δ 8.30 (1H), 7.74 (1H), 7.57 (2H), 7.48 (1H), 7.27 (1H), 5.29 (2H), 4.67–4.36 (2H), 4.53 (1H), 4.23 (1H), 3.37–3.11 (3H), 2.97 (1H), 1.27 (9H).

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Scheme I.



Scheme I teaches a general method for the preparation of azatyrosine reagents I-3, I-4, I-5, I-6 and I-7, by adaptation of the methodology for the preparation of protected azatyrosine reagents as described by Kawata, S.; Ashizawa, S.; Hirama, M. *J. Am. Chem. Soc.* 1997, 119, 12012-12013 and references cited therein. Thus *O*-protection of chloriodopyridinol I-1 (identical to H-2) is followed by reaction with the organozinc, derived from a suitably protected β -iodoalanine, to provide the protected 6-chloroazatyrosine I-3. Reductive dehalogenation of I-3 gives I-4, that is *O*-deprotected to give I-5. Reagent I-5 is *O*-alkylated, as exemplified by the preparation of I-6. *N*-deprotection of I-6 gives the aminoester I-7, that is used (as exemplified by reagent A-4 of Scheme A, and by reagent B-5 of Scheme B) for the synthesis of Examples of this invention.

Preparation 45.

(Scheme I, I-2)

(\pm)-2-Chloro-3-[(2-tetrahydropyranyl)oxy]-6-iodopyridine (C₁₀H₁₁ClINO₂).

To a solution of chloriodopyridinol I-1 (same as H-2, the product of preparation 41) (1.00 g, 3.91 mmol) and dihydropyran (1.0 mL, 10.6 mmol) in CH₂Cl₂ (10 mL) under Ar at rt is added pyridinium chloride (0.050 g). The reaction mixture is stirred for 72 h. It is diluted with CH₂Cl₂, and is washed with satd aq NaHCO₃ and brine. The CH₂Cl₂ solution is dried, filtered and concentrated to an oil, that is purified by silica flash chromatography (19:1 hexanes/EtOAc) to give 1.06 g of the title product: TLC (19:1 hexanes/EtOAc) *R_f* 0.24; ¹H NMR (CDCl₃, 300 MHz) δ 7.55 (1H), 7.17 (1H), 5.50 (1H), 3.77 (1H), 3.61 (1H), 2.07–1.57 (6H).

Preparation 46.

(Scheme I, I-3)

(*S*)-2-Chloro- α -[(1,1-dimethylethoxy)carbonyl]amino]-3-[(2-tetrahydropyranyl)oxy]-6-pyridinepropanoic acid methyl ester (C₁₉H₂₇ClN₂O₆).

To an amberized flask containing activated Zn dust (0.349 g, 5.51 mmol) under Ar is added THF (2 mL) and 1,2-dibromoethane (0.018 mL, 0.21 mmol). The suspension is

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brought to reflux for several minutes, cooled to approximately 30 °C, and TMSCl (0.17 mL of a 1 M solution in THF) is added. The reaction mixture is stirred at 40±5 °C for 30 min and then is cooled to below rt. A solution of *N*-[(1,1-dimethylethoxy)carbonyl]-3-iodo-L-alanine methyl ester [93267-04-0] (1.81 g, 5.50 mmol) in 11:7 dimethylacetamide/THF (9.0 mL) is added, and the resulting reaction mixture is stirred at 45 °C for 5 h. The reaction mixture is cooled to below rt, and solid PdCl₂(PPh₃)₂ (0.192 g) is added, followed immediately by addition of a degassed solution of the iodopyridine (0.936 g, 2.76 mmol) in 1:1 THF/dimethylacetamide (5.6 mL). This reaction mixture is stirred for 4 h at 45 °C. It is cooled to 0 °C, quenched with sat'd aq NH₄Cl, and extracted with EtOAc. The combined EtOAc portions are washed with sat'd aq NH₄Cl and brine. The EtOAc solution is dried, filtered and concentrated to give a green-yellow colored foam, that upon purification by silica flash chromatography (7:3 hexanes/EtOAc) gives 0.879 g (1.85 mmol, 60%) of the title product: TLC (7:3 hexanes/EtOAc) *R_f* 0.21; ¹H NMR (CDCl₃, 300 MHz) δ 7.39 (1H), 7.00 (1H), 5.46 (1H), 4.61 (1H), 4.13 (1H), 3.80 (3H), 3.62 (1H), 3.20 (1H), 2.13–1.53 (6H), 1.42 (9H).

Preparation 47.

(Scheme I, I-4)

(*S*)-α-[[[(1,1-Dimethylethoxy)carbonyl]amino]-3-[(2-tetrahydropyranyl)oxy]-6-pyridinepropanoic acid methyl ester (C₁₉H₂₈N₂O₆).

A suspension of pre-reduced Pd/CaCO₃ (3.5 g) and I-3 (1.15 g, 2.77 mmol) in EtOH (40 mL) is hydrogenated (30 psi H₂) for 19 h at rt. The mixture is filtered, and the filtrate is evaporated to give a yellow-colored foam, that is purified by silica flash chromatography (600:400:1 hexanes/EtOAc/*i*PrOH) to give 0.367 g of the title compound: TLC (1:1 hexanes/EtOAc) *R_f* 0.27; ¹H NMR (CDCl₃, 300 MHz) δ 8.30 (1H), 7.29 (1H), 7.03 (1H), 5.81 (1H), 5.39 (1H), 4.65 (1H), 3.86 (1H), 3.73 (3H), 3.62 (1H), 3.21 (2H), 1.96–1.53 (6H), 1.42 (9H).

Preparation 48.

(Scheme I, I-5)

(S)- α -[[1,1-Dimethylethoxy)carbonyl]amino]-5-hydroxy-2-pyridinepropanoic acid methyl ester (C₁₄H₂₀N₂O₅).

- 5 A solution of I-4 (0.346 g, 0.91 mmol) and pyridinium *p*-toluenesulfonate (0.031 g, 0.12 mmol) in EtOH (8 mL) is stirred at 55±5 °C for 20 h. The reaction mixture is cooled to rt, and concentrated *in vacuo*. The residue is taken up in EtOAc. The solution is washed with brine, and dried, filtered and concentrated to a pale yellow-colored oil that is purified by silica flash chromatography (500:500:1 hexanes/EtOAc/*i*PrOH). Evaporation of the
- 10 column fractions gives 0.132 g of the title compound: TLC (1:1 hexanes/EtOAc) *R_f* 0.18; ¹H NMR (CDCl₃, 300 MHz) δ 8.13 (1H), 7.13 (1H), 7.03 (1H), 5.71 (1H), 4.65 (1H), 3.70 (3H), 3.20 (2H), 1.39 (9H).

Preparation 49.

(Scheme I, I-6)

- 15 (S)-5-[(2,6-Dichlorophenyl)methoxy]- α -[[1,1-dimethylethoxy)carbonyl]amino]-2-pyridinepropanoic acid methyl ester (C₂₁H₂₄Cl₂N₂O₅).

- To a solution of I-5 (0.126 g, 0.43 mmol), 2,6-dichlorobenzylalcohol (0.075 g, 0.43 mmol) and PPh₃ (0.113 g, 0.43 mmol) in dry THF (4 mL) at 0 °C under Ar is added DEAD (0.068 mL). The reaction mixture is permitted to warm to rt, and is stirred for 18
- 20 h. It is concentrated, and the residue is purified by silica flash chromatography (700:300:1 hexanes/EtOAc/*i*PrOH) to give 0.149 g of the title compound: TLC (7:3 hexanes/EtOAc) *R_f* 0.34; ¹H NMR (CDCl₃, 300 MHz) δ 8.31 (1H), 7.37 (2H), 7.25 (2H), 7.08 (1H), 5.81 (1H), 5.29 (2H), 4.65 (1H), 3.70 (3H), 3.24 (2H), 1.63 (1 H), 1.43 (9H).

Preparation 50.

- 25 (Scheme I, I-7)

(S)- α -Amino-5-[(2,6-dichlorophenyl)methoxy]-2-pyridinepropanoic acid methyl ester dihydrogen chloride salt (C₁₆H₁₆Cl₂N₂O₃·2HCl).

A solution of carbamate I-6 (0.546 g, 1.20 mmol) in 4 M HCl in dioxane (12 mL) is stirred at rt under Ar for 16 h. The reaction mixture is concentrated *in vacuo*. The residue is dissolved in H₂O (40 mL), and this solution is extracted with Et₂O. The aqueous solution is frozen and lyophilized to give, as a light yellow-colored solid, 0.485 g of the
5 title compound: ¹H NMR (CD₃SOCD₃, 300 MHz) δ 8.75 (3H), 8.47 (1H), 7.81 (1H), 7.57 (3H), 7.48 (1H), 5.35 (2H), 4.49 (1H), 3.67 (3H), 3.42 (2H).

Example 120.

(Scheme A, A-5)

[S-(R*,R*)]-4-[[[1-[[3-[(2,6-Dichlorophenyl)methoxy]-6-pyridyl]methyl]-2-
10 methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid (1,1-dimethylethyl) ester (C₂₅H₃₀Cl₂N₃O₆S).

Example 120 was prepared as described in Scheme A from D-cysteine using the product of preparation 49 as amino acid intermediate A-4. Physical properties as follows: TLC (1:1 hexanes/EtOAc) *R_f* 0.22; ¹H NMR (CDCl₃, 300 MHz) δ 8.28 (2H), 7.38 (2H), 7.28 (2H),
15 7.09 (1H), 5.29 (2H), 4.90-4.74 (3H), 4.40 (1H), 3.67 (3H), 3.38-3.22 (3H), 1.61 (2H), 1.40 (9H).

Example 121.

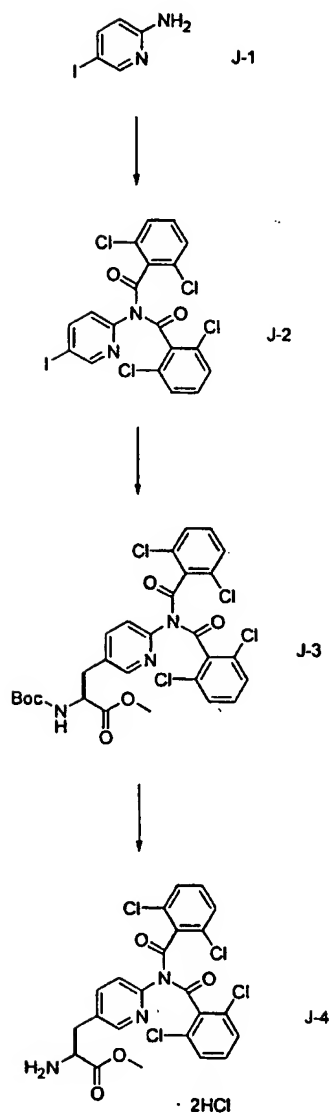
(Scheme A, A-9)

[S-(R*,R*)]-4-[[[1-Carboxy-2-[3-[(2,6-dichlorophenyl)methoxy]-6-
20 pyridyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid (1,1-dimethylethyl) ester (C₂₄H₂₈Cl₂N₃O₆S).

Example 121 was prepared from example 120 by the procedure described in preparation 12. Physical data as follows: mp 92-94°; TLC (500:500:3 hexanes/EtOAc/HCO₂H) *R_f* 0.10; ¹H NMR (CDCl₃, 300 MHz) δ 8.31 (1H), 8.26 (1H), 7.55 (2H), 7.28 (2H), 7.46
25 (1H), 7.21 (1H), 5.25 (2H), 4.72-4.38 (2H), 4.60 (1H), 4.23 (1H), 3.21-3.12 (2H), 3.09-2.94 (1H), 2.74 (1H), 1.29 (9H).

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Scheme J.



5

Scheme J teaches a general method for the preparation of *para*-acylamino derivatives of aza-phenylalanine. Thus bis-acylation of 2-amino-5-iodopyridine **J-1** gives the imide **J-2**, that is reacted with the organozinc, derived from a suitably protected β -iodoalanine, to provide the protected acylamino azaphenylalanine **J-3**. *N*-deprotection of **J-3** gives the

aminoester J-4, that is used (as exemplified by reagent A-4 of Scheme A, and by reagent B-5 of Scheme B) for the synthesis of Examples of this invention.

Preparation 51.

(Scheme J: J-2)

5 2-[[Bis(2,6-Dichlorobenzoyl)]amino]-5-iodopyridine (C₁₉H₉Cl₄IN₂O₂).

To a solution of 2-amino-5-iodopyridine J-1 (2.20 g, 10.0 mmol) and Et₃N (2.12 mL, 15.0 mmol) in dry THF (100 mL) at rt under Ar, is added dropwise 2,6-dichlorobenzoylchloride (1.60 mL, 11.0 mmol) over 45 min. The reaction mixture is stirred for 15 h. It is diluted with EtOAc (300 mL), and is washed with cold aq 1 M NaOH
10 and brine. The solution is dried, filtered and concentrated to give a yellow-colored waxy solid, that is purified by silica flash chromatography (3:1 hexanes/EtOAc) to give 2.60 g of the title compound: TLC (7:3 hexanes/EtOAc) *R_f* 0.60; ¹H NMR (CDCl₃, 300 MHz) δ 8.59 (1H), 8.03 (1H), 7.99 (1H), 7.44–7.26 (6H).

Preparation 52.

15 (Scheme J: J-3)

(S)-2-[[Bis-(2,6-dichlorobenzoyl)]amino]-α-[[[(1,1-dimethylethoxy)carbonyl]amino]-5-pyridinepropanoic acid methyl ester (C₂₈H₂₅Cl₄N₃O₆).

To an amberized flask containing activated Zn dust (0.865 g, 13.23 mmol) under Ar is added sequentially *N*-[(1,1-dimethylethoxy)carbonyl]-3-iodo-L-alanine methyl ester
20 (4.36 g, 13.23 mmol), THF (13 mL) and *N,N*-dimethylacetamide (13 mL). The reaction mixture is purged of O₂ by the bubbling Ar through the mixture for 5 min, and then is warmed to 45±5 °C for 7 h. It is cooled to rt. To this mixture is added PdCl₂(PPh₃)₂ (0.461 g) followed immediately by a degassed solution of iodide J-2 (2.60 g, 4.59 mmol)
25 in 1:1 THF/*N,N*-dimethylacetamide (18 mL). The reaction mixture is stirred at 45±5 °C under Ar for 13 h. It is cooled to 0 °C and quenched with satd aq NH₄Cl (150 mL). The mixture is extracted with EtOAc. The combined EtOAc extracts are washed with brine, dried, filtered and concentrated to a green-yellow-colored paste, that is purified by silica

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flash chromatography (700:300:1 hexanes/EtOAc/iPrOH) to give 1.43 g of the title compound: TLC (7:3 hexanes/EtOAc) R_f 0.29; ^1H NMR (CDCl_3 , 300 MHz) δ 8.13 (1H), 7.57 (1H), 7.46 (1H), 7.26 (6H), 4.80 (1H), 4.50 (1H), 3.67 (3H), 3.05 (2H), 1.46 (9H).

Preparation 53.

5 (Scheme J: J-4)

(S)- α -Amino-2-[[bis-(2,6-dichlorobenzoyl)]amino]-5-pyridinepropanoic acid methyl ester dihydrochloride salt ($\text{C}_{23}\text{H}_{17}\text{Cl}_4\text{N}_3\text{O}_4\cdot\text{HCl}$).

A solution of J-3 (0.69 g, 1.08 mmol) in 4 M HCl in dioxane (15 mL) is stirred under Ar for 20 h. The reaction mixture is concentrated *in vacuo*, diluted with H_2O , and extracted
10 with Et_2O . The aqueous solution is frozen and lyophilized to give, as a pale yellow-colored solid, 0.627 g of the title compound: ^1H NMR (CD_3SOCD_3 , 300 MHz) δ 8.80 (2H), 8.27 (1H), 7.82 (1H), 7.67–7.26 (7H), 4.25 (1H), 3.52 (3H), 3.16 (2H), 3.04 (1H); MS (ESI+) m/z 541.7.

Example 122.

15 (Scheme A, A-5)

[S-(R*,R*)]-4-[[[1-[[2-[[Bis-(2,6-dichlorobenzoyl)]amino]-5-pyridyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid (1,1-dimethylethyl) ester ($\text{C}_{25}\text{H}_{30}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$).

Example 122 was prepared as described in Scheme A from D-cysteine using the product of
20 preparation 52 as amino acid intermediate A-4. Physical properties as follows: TLC (1:1 hexanes/EtOAc) R_f 0.22; ^1H NMR (CDCl_3 , 300 MHz) δ 8.28 (2H), 7.38 (2H), 7.28 (2H), 7.09 (1H), 5.29 (2H), 4.90–4.74 (3H), 4.40 (1H), 3.67 (3H), 3.38–3.22 (3H), 1.61 (2H), 1.40 (9H).

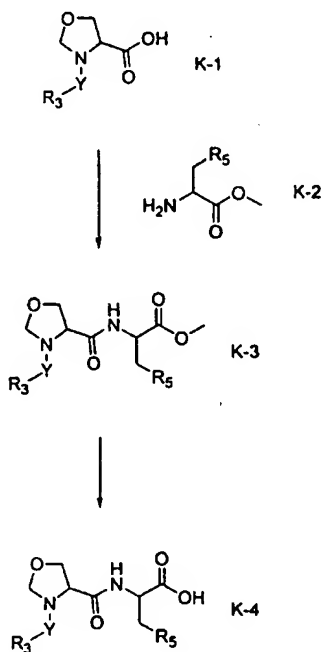
Example 123.

25 (Scheme A, A-9)

[S-(R*,R*)]-4-[[[1-Carboxy-2-[2-[[Bis-(2,6-dichlorobenzoyl)]amino]-5-pyridyl]]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid (1,1-dimethylethyl) ester ($\text{C}_{31}\text{H}_{28}\text{Cl}_4\text{N}_4\text{O}_7\text{S}$).

Example 123 was prepared from example 122 by the procedure described in preparation 12. Physical data as follows: mp 158–160°; TLC (50:50:2 hexanes/EtOAc/HCO₂H) *R_f* 0.18; ¹H NMR (CD₃SOCD₃, 300 MHz) δ 8.42 (1H), 8.22 (1H), 7.75 (1H), 7.69–7.17 (5H), 7.53 (1H), 4.59 (1H), 4.40 (2H), 4.19 (1H), 3.19–3.02 (2H), 2.84 (1H), 2.56 (1H), 1.34 (9H).

Scheme K.



10

Scheme K teaches a general method for the preparation of oxazolidinecarboxylic acid Examples K-3 and K-4, where R₃, R₅ and Y are identical to the definitions of Scheme B. Thus coupling of oxazolidinecarboxylic acid K-1 and aminoester K-2 (as exemplified by the reaction of reagents A-3 and A-4 of Scheme A, and B-4 and B-5 of Scheme B) provides Examples K-3, that are hydrolyzed to Examples K-4 of this invention.

Example 124.

(Scheme K: K-3, where R₃ is (phenyl)methyl, R₂ is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is CO₂ and the stereochemistry is [S-(R*,R*)])

5 [S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-oxazolidinecarboxylic acid 3-phenylmethyl ester (C₂₉H₂₇Cl₂N₃O₇).

Example 124 was prepared as by coupling commercially available (S)-3,4-oxazolidinedicarboxylic acid 3-(phenylmethyl) ester to amino acid K-2 (Scheme K, where R₃ is 4-[(2,6-dichlorobenzoyl)amino]phenyl stereochemistry is (S)) under the conditions described by
 10 preparation 3. Physical properties as follows: TLC (3:2 Heptane/EtOAc) R_f= 0.17; UV (MeOH) λ_{max} 225 (ε 12600, sh), 251 (17900); ¹³C NMR (d₆-dimethylsulfoxide) δ 188.75, 171.96, 169.86, 162.40, 162.30, 152.99, 137.59, 137.47, 136.87, 136.81, 133.23, 131.79, 131.65, 130.08, 128.80, 128.67, 128.28, 127.76, 119.84, 119.75, 79.82, 66.77, 57.85, 53.85, 52.47, 36.62 (23 lines expected; 26 lines observed); MS (FAB) m/z 602, 600, 558,
 15 556, 531, 466, 371, 351, 349, 280, 278, 175, 173; MS (FAB) m/z 600.1312 (calcd [M+H]⁺ 600.1304; Anal. C, 57.75; H, 4.75; N, 6.80; Cl, 11.86 (calcd for 0.42% H₂O: C 57.77, H 4.56, N 6.97, Cl 11.76).

Example 125.

(Scheme K: K-4 where R₃ is (phenyl)methyl, R₂ is 4-[(2,6-dichlorobenzoyl)amino]phenyl,
 20 Y is CO₂ and the stereochemistry is [S-(R*,R*)])

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]-carbonyl]-3-oxazolidinecarboxylic acid 3-phenylmethyl ester (C₂₈H₂₅Cl₂N₃O₇).

Example 125 was prepared from example 124 by the procedure described in preparation 12.

25 Physical properties as follows: TLC (950:50:1 CH₂Cl₂/MeOH/HCO₂H) R_f= 0.34; ¹H NMR (d₆-dimethylsulfoxide) δ 10.65 (1H), 8.31 (1H), 7.61-7.42 (5H), 7.40-7.20 (5H), 7.15 (2H), 5.19-4.85 (2H), 4.90 (1H), 4.76 (1H), 4.43 (1H), 4.36 (1H), 4.11 (1H), 3.65 (1H), 3.04 (1H), 2.87 (1H); MS (FAB) m/z 588, 586 544, 542, 532, 391, 337, 335, 327, 269, 267, 161, 147.133, 129, 117, 115, 103, 101, 91; MS (FAB) 586.1132 (calcd
 30 586.1147).

Example 126.

(Scheme K: K-3 where R_3 is benzyl, R_4 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is CO_2 and the stereochemistry is $[R-(R^*,S^*)]$)

$[R-(R^*,S^*)]$ -4-[[[1-[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-oxazolidinecarboxylic acid 3-phenylmethyl ester
($\text{C}_{29}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7$).

Example 126 was prepared as described in Scheme K from commercially available (*R*)-3,4-oxazolidinedicarboxylic acid 3-(phenylmethyl) ester. Physical properties as follows: TLC (3:2 heptane/EtOAc) R_f = 0.19; UV (MeOH) λ_{max} 225 (e 12400, sh), 252 (17700), 284 (2960, sh); ^{13}C NMR (CDCl_3) δ 171.23, 169.14, 162.43, 154.26, 136.47, 135.93, 135.65, 132.37, 132.28, 130.91, 130.00, 128.62, 128.39, 128.11, 120.30, 79.66, 67.94, 58.38, 53.14, 52.49, 37.19 (23 lines expected; 21 lines observed); MS (FAB) m/z 602, 600, 558, 556, 466, 351, 349, 280, 278, 175, 173; MS (FAB) m/z 600.1299 (calcd for $[\text{M}+\text{H}]^+$ 600.1304); Anal. C 57.69, H 4.90, N 6.71, Cl 11.49 (calcd for 0.35% H_2O : C 57.81, H 4.56, N 6.97, Cl 11.77).

Example 127.

(Scheme K: K-4 where R_3 is benzyl, R_4 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is CO_2 and the stereochemistry is $[R-(R^*,S^*)]$)

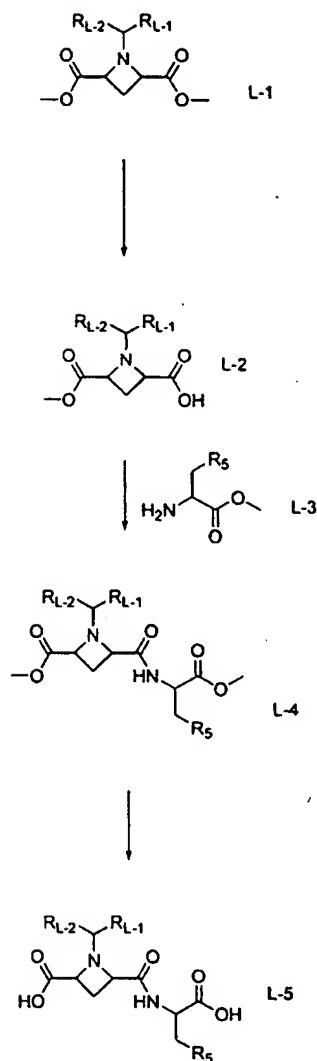
$[R-(R^*,S^*)]$ -4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]-carbonyl]-3-oxazolidinecarboxylic acid 3-(phenylmethyl) ester ($\text{C}_{28}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_7$).

Example 127 was prepared from example 126 by the procedure described in preparation 12.

Physical properties as follows: TLC (950:50:1 $\text{CH}_2\text{Cl}_2/\text{MeOH}/\text{HCO}_2\text{H}$) R_f = 0.31; ^1H NMR (d_4 -methanol) δ 7.60 (2H), 7.49-7.23 (8H), 7.21 (2H), 5.23-4.97 (2H), 4.95 (1H), 4.88 (1H), 4.70 (1H), 4.36 (1H), 4.14 (1H), 3.85-3.73 (1H), 3.23 (1H), 3.00 (1H); MS (FAB) m/z 588, 586, 544, 542, 532, 391, 371, 337, 335, 245, 177, 173, 149, 123, 105, 103, 91; MS (FAB) m/z 586.1163 (calcd 586.1147).

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Scheme L.



- 5 Scheme L teaches a general method for the preparation of *N*-alkylaryl azetidinedi-carboxylic acid Examples L-4 and L-5, where R_{L-1} is C_{1-6} alkyl, R_{L-2} is C_{6-10} aryl and R_5 is defined as in Scheme B. The *N*-phenylethyl-2,4-azetidinedicarboxylic acid dimethyl ester stereoisomers of general structure L1 were prepared as described (Hoshino, J.; Hiraoka, J.; Hata, Y.; Sawada, S.; Yamamoto, Y. *J. Chem. Soc. Perkin Trans. I* 1995, 693-697) and
- 10 separated by silica flash chromatography. Thus partial saponification of diester L-1 gives

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the half-acid L-2, that is coupled with reagent L-3 (as exemplified by the use of reagents A-3 and A-4 of Scheme A, and B-4 and B-5 of Scheme B) to provide Examples L-4, that are then hydrolyzed to Examples L-5 of this invention.

Preparation 54.

- 5 (Scheme L: L-1 where R_{L-1} is methyl, R_{L-2} is phenyl and stereochemistry is [2*S*-[1(*R*^{*}),2 α ,4 β]]).
- [2*S*-[1(*R*^{*}),2 α ,4 β]]-1-(1-Phenylethyl)-2,4-azetidinedicarboxylic acid dimethyl ester**
[168647-92-5] ($C_{15}H_{19}NO_4$).

Physical properties as follows: TLC (4:1 Hexanes/EtOAc) R_f = 0.42; ^{13}C NMR ($CDCl_3$) δ
10 173.61, 142.41, 128.31, 127.39, 127.35, 60.95, 60.67, 51.64, 24.96, 21.72.

Preparation 55.

- (Scheme L: L-1 where R_{L-1} is methyl, R_{L-2} is phenyl and stereochemistry is [2*R*-[1(*S*^{*}),2 α ,4 β]]).
- [2*R*-[1(*S*^{*}),2 α ,4 β]]-1-(1-Phenylethyl)-2,4-azetidinedicarboxylic acid dimethyl ester**
15 **($C_{15}H_{19}NO_4$).**

Physical properties as follows: TLC (4:1 Hexanes/EtOAc) R_f = 0.31; ^{13}C NMR ($CDCl_3$) δ
172.97, 141.21, 128.44, 128.07, 127.68, 61.54, 60.69, 51.61, 24.91, 19.55.

Preparation 56.

- (Scheme L: L-1 where R_{L-1} is methyl, R_{L-2} is phenyl and stereochemistry is [1(*S*)-*cis*]).
- 20 **[1(*S*)-*cis*]-1-(1-Phenylethyl)-2,4-azetidinedicarboxylic acid dimethyl ester [168753-32-0] ($C_{15}H_{19}NO_4$).**

Physical properties as follows: TLC (4:1 Hexanes/EtOAc) R_f = 0.21; ^{13}C NMR ($CDCl_3$) δ
172.58, 172.08, 140.84, 128.22, 128.15, 127.65, 66.32, 60.10, 59.65, 52.06, 51.59, 24.26,
19.91.

- 25 Preparation 57.

(Scheme L: L-1 where R_{L-1} is methyl, R_{L-2} is phenyl and stereochemistry is [2*R*-[1(*R*^{*}),2 α ,4 β]]).

[2*R*-[1(*R*^{*}),2 α ,4 β]]-1-(1-Phenylethyl)-2,4-azetidinedicarboxylic acid dimethyl ester
($C_{15}H_{19}NO_4$).

Physical properties as follows: TLC (8:2 hexanes/EtOAc) R_f = 0.42; ^{13}C NMR (CDCl_3) δ 173.61, 142.41, 128.31, 127.39, 127.35, 60.95, 60.67, 51.64, 24.96, 21.72; MS (+ESI) m/z 278.3.

Preparation 58.

- 5 (Scheme L: L-1 where $R_{L,1}$ is methyl, $R_{L,2}$ is phenyl and stereochemistry is [2*S*-[1(*S*^{*}),2 α ,4 β]].
[2*S*-[1(*S*^{*}),2 α ,4 β]]-1-(1-Phenylethyl)-2,4-azetidinedicarboxylic acid dimethyl ester
($\text{C}_{15}\text{H}_{19}\text{NO}_4$).

Physical properties as follows: TLC (8:2 hexanes/EtOAc) R_f = 0.31; ^{13}C NMR (CDCl_3) δ
10 172.97, 141.21, 128.44, 128.07, 127.68, 61.54, 60.69, 51.61, 24.91, 19.55; MS (+ESI) m/z 278.3.

Preparation 59.

- (Scheme L: L-1 where $R_{L,1}$ is methyl, $R_{L,2}$ is phenyl and stereochemistry is [1(*R*)-*cis*]].
[1(*R*)-*cis*]-1-(1-Phenylethyl)-2,4-azetidinedicarboxylic acid dimethyl ester
15 ($\text{C}_{15}\text{H}_{19}\text{NO}_4$).
Physical properties as follows: TLC (8:2 hexanes/EtOAc) R_f = 0.21; ^{13}C NMR (CDCl_3) δ
172.58, 172.08, 140.84, 128.22, 128.15, 127.65, 66.32, 60.10, 59.65, 52.06, 51.59, 24.26,
19.91; MS (+ESI) m/z 278.3.

Preparation 60.

- 20 (Scheme L: L-2 where $R_{L,1}$ is methyl, $R_{L,2}$ is phenyl and stereochemistry is [2*S*-[1(*R*^{*}),2 α ,4 β]]).
[2*S*-[1(*R*^{*}),2 α ,4 β]]-1-(1-Phenylethyl)-2,4-azetidinedicarboxylic acid monomethyl ester
($\text{C}_{14}\text{H}_{17}\text{NO}_4$).
A mixture of L-1 (Scheme L, where $R_{L,1}$ is methyl, $R_{L,2}$ is phenyl and stereochemistry is
25 [2*S*-[1(*R*^{*}),2 α ,4 β]], the product of preparation 54) (7.95 g; 28.7 mmol) and LiOH (30
mmol) in 1:1 MeOH/ H_2O (240 mL) is stirred at rt for 42 h. The reaction mixture is
adjusted to pH 5 with HOAc, and is concentrated. The resulting concentrate is diluted with
brine and extracted repeatedly with CHCl_3 . The combined CHCl_3 extracts are dried,
filtered and concentrated to give a yellow foam (6.61 g), that is purified by preparative
30 C18 reverse phase chromatography to give the title compound as a crystalline solid: mp

112-113 °C; TLC (650:350:1 hexanes/EtOAc/HCO₂H) R_f = 0.17; MS (FAB) m/z 527, 264, 248, 218, 204, 192, 186, 177, 160, 114, 105; Anal. C 64.04, H 6.57, N 5.37 (calcd C 63.87, H 6.51, N 5.32).

Preparation 61 and Example 128.

- 5 (Scheme L: L-4 where R_{L-1} is methyl, R_{L-2} is phenyl, R_3 is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [2*S*-[1(*R**),2 α ,4 β (*R**)]])
- [2*S*-[1(*R**),2 α ,4 β (*R**)]]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid methyl ester (C₃₁H₃₁Cl₂N₃O₆).
- 10 A mixture of L-2 (Scheme L, where R_{L-1} is methyl, R_{L-2} is phenyl and stereochemistry is [2*S*-[1(*R**),2 α ,4 β]], the product of preparation 60) (0.62 g, 2.4 mmol), L-3 (Scheme L where R_3 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*)) (0.95 g, 2.4 mmole), and BOP-Cl (0.68 g; 2.7 mmol) in CH₂Cl₂ (10 mL) is treated with (*i*-Pr)₂NEt (1.7 mL, 9.8 mmol). The reaction mixture is stirred at rt under N₂ for 19 h. It is diluted with
- 15 half-saturated NaHCO₃ and extracted with CH₂Cl₂. The CH₂Cl₂ extracts are dried, filtered and concentrated to give a beige-colored foam (1.40 g), that is purified by silica flash chromatography to give the title compound: TLC (1:1 Hexanes/EtOAc) R_f = 0.23; ¹³C NMR (CDCl₃) δ 173.25, 172.73, 171.63, 162.57, 141.86, 136.76, 136.01, 132.44, 132.39, 130.90, 130.00, 128.57, 128.12, 127.55, 127.08, 120.37, 63.25, 60.06, 59.17, 52.41, 52.12,
- 20 51.47, 37.65, 26.06, 21.25; MS (FAB) m/z 612.1658; Anal. C 59.66, H 5.14, N 6.64 (calcd C 60.79, H 5.10, N, 6.86).

Preparation 62 and Example 129.

- (Scheme L: L-5 where R_{L-1} is methyl, R_{L-2} is phenyl, R_3 is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [2*S*-[1(*R**),2 α ,4 β (*R**)]])
- 25 [2*S*-[1(*R**),2 α ,4 β (*R**)]]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]-ethyl]-amino]carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid dilithium salt (C₂₉H₂₅Cl₂Li₂N₃O₆).
- A mixture of L-4 (Scheme L, where R_{L-1} is methyl, R_{L-2} is phenyl, R_3 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is [2*S*-[1(*R**),2 α ,4 β (*R**)]]], the product
- 30 of preparation 61) (0.684 g, 1.12 mmol) and LiOH·H₂O (0.25 g, 6.0 mmol) is dissolved by

warming in MeOH (10 mL). This solution is diluted with 1:1 H₂O/THF (20 mL), and the reaction mixture is stirred at rt for 22 h. The mixture is adjusted with 1 N HCl to pH 6. The solution is concentrated to give a white solid, that is purified by preparative C18 reverse phase chromatography (MeCN/H₂O gradient). Evaporation of the column fractions gives a white solid, that is dissolved in warm H₂O. The solution is frozen and lyophilized to give, as a white solid, the title compound: mp 270 °C; TLC (850:150:1 CHCl₃/MeOH/HCO₂H) *R_f* = 0.21-0.36; ¹³C NMR (CD₃OD) δ 178.05, 174.59, 174.01, 162.20, 142.33, 135.09, 134.91, 133.04, 130.54, 129.24, 128.61, 126.46, 126.41, 126.17, 124.94, 118.33, 82.70, 61.70, 60.50, 53.30, 36.23, 25.85, 18.65; MS (FAB) *m/z* 584.1350; Anal. C 55.17, H 5.01, N 6.63, Cl 11.16 (calcd for 7.52% H₂O: C 54.02, H 4.75, N 6.52, Cl 11.21).

Example 130.

(Scheme L: L-4 where R_{L-1} is methyl, R_{L-2} is phenyl, R₃ is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [2*R*-[1(*S**),2α,4β(*S**)]]).

[2*R*-[1(*S**),2α,4β(*S**)]]-4-[[[1-[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid methyl ester (C₃₁H₃₁Cl₂N₃O₆).

Example 130 was prepared as described in Scheme L using the product of preparation 55 as intermediate L-1. Physical properties as follows: TLC (3:2 EtOAc/Hexanes) *R_f* = 0.43; [α]_D²⁵ +58 (c 0.91, MeOH); ¹³C NMR (CDCl₃) δ 173.24, 172.15, 172.05, 162.58, 140.75, 136.45, 136.01, 132.89, 132.38, 130.91, 129.60, 128.67, 128.39, 128.11, 127.97, 120.39, 63.54, 60.19, 59.54, 53.15, 52.16, 51.91, 37.86, 26.12, 18.37. MS (EI) *m/z* 613, 611, 598, 596, 554, 552, 527, 525, 508, 506, 450, 448, 351, 349, 218, 191, 175, 173, 160, 131, 114, 105; Anal. C 60.68, H 5.18, N 6.67, Cl 11.24 (calcd for C 60.79, H 5.10, N 6.86, Cl 11.58).

Example 131.

(Scheme L: L-5 where R_{L-1} is methyl, R_{L-2} is phenyl, R₃ is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [2*R*-[1(*S**),2α,4β(*S**)]]).

[2*R*-[1(*S**),2α,4β(*S**)]]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)-amino]phenyl]ethyl]-amino]carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid, dilithium salt (C₂₉H₂₅Cl₂Li₂N₃O₆).

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Example 131 was prepared from Example 130 by the procedure described in preparation 62. Physical properties as follows: $[\alpha]_D^{25} +84$ (c 0.95, MeOH); ^{13}C NMR (CD_3OD) δ 179.18, 176.55, 174.13, 163.58, 141.88, 136.32, 136.29, 134.96, 131.90, 130.86, 129.47, 128.49, 128.07, 127.87, 127.16, 119.74, 63.70, 63.49, 59.77, 56.27, 38.24, 26.25, 18.70; MS (FAB) m/z 598, 596, 592, 590, 552, 550, 546, 544, 161; Anal. C 55.20, H 5.52, N, 6.59 (calcd for 10.98% H_2O : C 52.00, H 4.99, N 6.27).

Example 132.

(Scheme L: L-4 where $\text{R}_{\text{L-1}}$ is methyl, $\text{R}_{\text{L-2}}$ is phenyl, R_5 is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [1(*S*),2 α ,4 α (*S*)], a single diastereomer having a *cis* relative configuration but unknown absolute configuration at C-2 and C-4 of the azetidine.)

[1(*S*),2 α ,4 α (*S*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid 2-methyl ester ($\text{C}_{31}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6$).

Example 132 was prepared as described in Scheme L using the product of preparation 56 as intermediate L-1. Physical properties as follows: TLC (9:1 CHCl_3 /acetone) $R_f = 0.29$; ^{13}C NMR (CDCl_3) δ 172.70, 172.29, 171.65, 162.49, 140.97, 136.71, 136.12, 132.47, 132.39, 130.81, 130.22, 128.35, 128.07, 127.82, 120.17, 66.61, 61.52, 59.99, 52.38, 52.24, 51.79, 37.73, 25.16, 20.09.

Example 133.

(Scheme L: L-5 where $\text{R}_{\text{L-1}}$ is methyl, $\text{R}_{\text{L-2}}$ is phenyl, R_5 is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [1(*S*),2 α ,4 α], a single diastereomer having a *cis* relative configuration but unknown absolute configuration at C-2 and C-4 of the azetidine.)

[1(*S*),2 α ,4 α (*S*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]-amino]-carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid ($\text{C}_{29}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_6$).

Example 133 was prepared from Example 132 by the procedure described in preparation 62. Physical properties as follows: $[\alpha]_D^{25} +20$ (c 0.88, MeOH); ^{13}C NMR (CD_3OD) δ 173.70, 173.05, 172.61, 163.69, 140.09, 136.93, 136.20, 133.18, 131.89, 130.89, 129.95, 128.09, 127.89, 127.88, 127.65, 120.03, 66.07, 61.29, 59.86, 52.70, 36.98, 25.03, 18.81;

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MS (FAB) m/z 662, 660, 586, 584, 539, 482, 480, 436, 434, 204, 175, 173, 160, 133, 109, 105; Anal. C 57.90, H 5.01, N 6.93 (calcd for 4.98% H₂O: C 56.63, H 4.98, N 6.83).

Example 134.

(Scheme L: L-4 where R_{L-1} is methyl, R_{L-2} is phenyl, R₅ is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [2*R*-[1(*R**),2 α ,4 β (*S**)]]]
 5 [2*R*-[1(*R**),2 α ,4 β (*S**)]]-4-[[[1-[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid methyl ester (C₃₁H₃₁Cl₂N₃O₆).

Example 134 was prepared as described in Scheme L using the product of preparation 57
 10 as intermediate L-1. Physical properties as follows: TLC (1:1 EtOAc/Hexanes) R_f =0.30; ¹³C NMR (CDCl₃) δ 173.78, 172.76, 172.10, 162.47, 141.81, 136.70, 135.96, 132.43, 132.38, 130.88, 129.79, 128.55, 128.10, 127.58, 127.00, 120.58, 63.32, 60.06, 59.27, 52.70, 52.50, 51.44, 37.08, 25.76, 21.52; MS (+ESI, 200:1 MeOH/HCO₂H solution) m/z 613.8, 611.8.

15 Example 135.

(Scheme L: L-5 where R_{L-1} is methyl, R_{L-2} is phenyl, R₅ is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [2*R*-[1(*R**),2 α ,4 β (*S**)]]]
 [2*R*-[1(*R**),2 α ,4 β (*S**)]]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]-ethyl]-amino]carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid dilithium salt
 20 (C₂₉H₂₅Cl₂Li₂N₃O₆).

Example 135 was prepared from Example 134 by the procedure described in preparation 62. Physical properties as follows: [α]_D²⁵ +76 (c 0.89, MeOH); ¹³C NMR (CD₃OD) δ 179.44, 176.71, 175.90, 163.55, 143.81, 136.54, 136.26, 134.47, 131.90, 120.81, 129.70, 127.83, 127.34, 126.33, 119.88, 63.06, 62.11, 58.71, 55.46, 37.17, 26.76, 20.56; MS
 25 (FAB) m/z 598, 596, 592, 590, 552, 550, 546, 544, 237, 105; Anal. C 55.12, H 5.24, N 6.59, Cl 10.56 (calcd for 6.21% H₂O: C 54.78, H 4.66, N 6.61, Cl 11.15).

Example 136.

(Scheme L: L-4 where R_{L-1} is methyl, R_{L-2} is phenyl, R₅ is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [2*S*-[1(*S**),2 α ,4 β (*R**)]]])

[2*S*-[1(*S*^{*}),2 α ,4 β (*R*^{*})]-4-[[[1-[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid methyl ester (C₃₁H₃₁Cl₂N₃O₆)

Example 136 was prepared as described in Scheme L using the product of preparation 58 as intermediate L-1. Physical properties as follows: TLC (85:15 CHCl₃/acetone) *R_f*= 0.54; ¹³C NMR (CDCl₃) δ 173.44, 172.62, 171.72, 162.79, 140.46, 136.69, 136.34, 133.12, 132.79, 131.29, 130.40, 129.08, 128.52, 128.45, 128.23, 120.57, 64.15, 60.51, 60.09, 53.11, 52.59, 52.19, 37.73, 26.70, 18.75; Anal. C 60.70, H 5.39, N 6.62 (calcd C 60.79, H 5.10, N 6.86).

10

Example 137.

(Scheme L: L-5 where R_{L-1} is methyl, R_{L-2} is phenyl, R₅ is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [2*S*-[1(*S*^{*}),2 α ,4 β (*R*^{*})]])

[2*S*-[1(*S*^{*}),2 α ,4 β (*R*^{*})]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]-ethyl]-amino]carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid (C₂₉H₂₇Cl₂N₃O₆).

15

Example 137 was prepared from Example 136 by the procedure described in preparation 62. Physical properties as follows: [α]_D²⁵ +2 (*c* 1.00, MeOH); ¹³C NMR (CD₃OD) δ 172.31, 169.43, 167.37, 163.72, 161.39, 160.93, 136.98, 136.11, 134.40, 133.20, 131.84, 130.93, 129.42, 129.39, 128.60, 127.89, 120.11, 62.68, 61.41, 53.51, 36.35, 25.00, 16.17; MS (FAB) *m/z* 586, 584, 482, 480, 204, 175, 173, 106, 105; Anal. C 53.07, H 4.52, N 6.15, Cl 10.46 (calcd for 0.80 equiv TFA and 2.13% H₂O: C 53.20, H 4.30, N 6.08, Cl 10.26).

20

Example 138.

(Scheme L: L-4 where R_{L-1} is methyl, R_{L-2} is phenyl, R₅ is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [1(*R*),2 α ,4 α (*S*)], a single diastereomer having a *cis* relative configuration but unknown absolute configuration at C-2 and C-4 of the azetidine.)

25

[1(*R*),2 α ,4 α (*S*)]-4-[[[1-[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid 2-methyl ester

30

(C₃₁H₃₁Cl₂N₃O₆).

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Example 138 was prepared as described in Scheme L using the product of preparation 59 as intermediate L-1. Physical properties as follows: TLC R_f = 0.47 (85:15 CHCl_3 /acetone); ^{13}C NMR (CDCl_3) δ 173.42, 172.40, 171.74, 162.46, 140.80, 136.76, 136.03, 132.64, 132.35, 130.80, 129.82, 128.38, 128.04, 127.85, 127.69, 120.39, 66.11, 61.50, 59.78, 52.83, 52.42, 51.72, 37.12, 25.03, 20.20; MS (+ESI, 200:1 MeOH/ HCO_2H solution) m/z 614.2, 612.2; Anal. C 60.66, H 5.18, N 6.80, Cl 11.42 (calcd C 60.79, H 5.10, N 6.86, Cl 11.58).

Example 139.

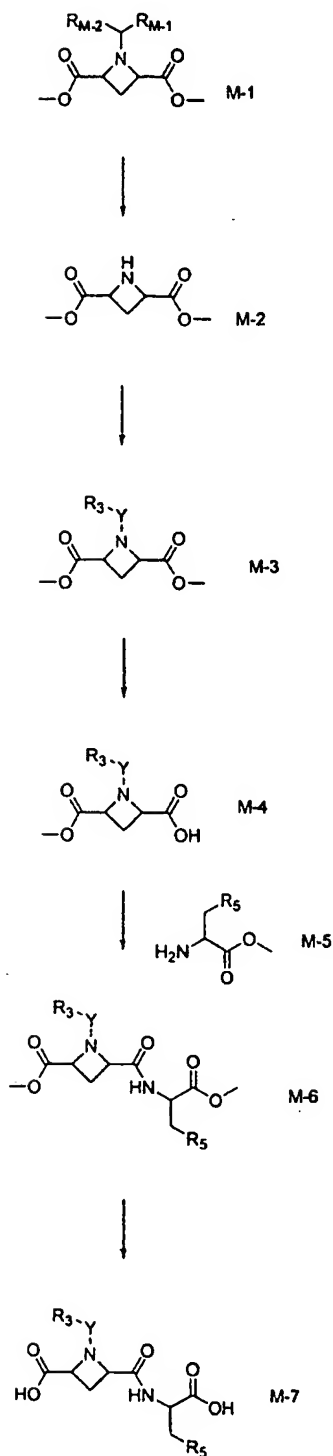
(Scheme L: L-5 where R_{L-1} is methyl, R_{L-2} is phenyl, R_1 is 4-[(2,6-dichlorobenzoyl)-amino]phenyl and stereochemistry is [1(R),2 α ,4 α (S)], a single diastereomer having a *cis* relative configuration but unknown absolute configuration at C-2 and C-4 of the azetidine)

[1(R),2 α ,4 α (S)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]-amino]-carbonyl]-1-(1-phenylethyl)-2-azetidinecarboxylic acid ($\text{C}_{29}\text{H}_{25}\text{Cl}_2\text{N}_3\text{O}_6$).

Example 139 was prepared from Example 138 by the procedure described in preparation 62. Physical properties as follows: $[\alpha]_D^{25}$ -30 (c 0.90, MeOH); ^{13}C NMR (CD_3OD) δ 172.71, 172.44, 171.81, 163.71, 138.03, 136.99, 136.15, 133.31, 131.89, 130.89, 129.45, 128.33, 128.29, 128.02, 127.86, 120.23, 65.62, 61.51, 59.60, 53.47, 36.04, 24.91, 17.68; MS (FAB) m/z 586, 584, 371, 298, 204, 177, 175, 173, 133, 105, 100; Anal. C 55.69, H 4.49, N 6.55, Cl 11.81 (calcd for 0.38 equiv TFA and 2.06% H_2O : C 55.80, H 4.54, N 6.56, Cl 11.07).

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Scheme M.



Scheme M teaches a general method for the preparation of *N*-acyl azetidinedicarboxylic acid Examples M-6 and M-7, where R_{M-1} is C_{1-6} alkyl, R_{M-2} is C_{6-10} aryl, and R_3 , R_5 and Y are defined as in Scheme B. Thus removal of the *N*-alkylaryl substituents R_{M-1} and R_{M-2} of M-1 gives aminodiester M-2, that is acylated to provide M-3. Partial saponification of diester M-3 gives half-acid M-4, that is coupled with reagent M-5 (as exemplified by the use of reagents A-3 and A-4 of Scheme A, and B-4 and B-5 of Scheme B) to provide Examples M-6, that are then hydrolyzed to Examples M-7 of this invention.

Preparation 63.

(Scheme M: M-2 where stereochemistry is (2*R*-*trans*))

10 (2*R*-*trans*)-2,4-Azetidinedicarboxylic acid dimethyl diester ($C_7H_{11}NO_4$).

To an N_2 -purged solution of freshly chromatographed Preparation M-1 (Scheme M where R_{M-1} is methyl, R_{M-2} is phenyl and stereochemistry is [2*R*-[1(*R**),2 α ,4 β]] prepared as described by preparation 57) (656 mg, 2.37 mmol) in MeOH (20 mL) is added 20% Pd(OH)₂/C (120 mg), and this mixture is hydrogenated for 19 h under an H_2 atmosphere (approximately 42 psi pressure). The mixture is filtered and concentrated to give the title compound as a colorless oil: TLC (4:1 Hexanes/EtOAc) R_f = 0.04; 1H NMR ($CDCl_3$) δ 4.33 (2H), 3.77 (6H), 3.38 (1H), 2.71 (2H); MS (+ESI) m/z 174.2.

Preparation 64.

(Scheme M: M-3 where R_3 is ethyl, Y is CO_2^- , and the stereochemistry is (2*R*-*trans*))

20 (2*R*-*trans*)-1,2,4-Azetidinetricarboxylic acid 1-ethyl-2,4-dimethyl triester ($C_{10}H_{15}NO_6$).

To a mixture of freshly prepared amine M-2 (Scheme M where stereochemistry is (2*R*-*trans*)) (14 mmol) in CH_2Cl_2 (20 mL) under N_2 at 0 °C is added Et_3N (3.0 mL, 22 mmol), followed by the dropwise addition of $ClCO_2Et$ (1.5 mL, 18 mmol). After 22 h the reaction is quenched with saturated $NaHCO_3$, diluted with H_2O , and extracted with EtOAc. The combined organic extracts are dried, filtered and concentrated to give 2.79 g of the carbamate, that is purified by silica flash chromatography: TLC (4:1 Hexanes/EtOAc) R_f = 0.17; $[\alpha]_D^{25} +183$ (c 0.83, MeOH); 1H NMR ($CDCl_3$) δ 4.77 (2H), 4.15 (1H), 4.10 (1H), 3.80 (6H), 2.58 (2H), 1.23 (3H); ^{13}C NMR (CD_3OD) δ 173.11, 173.05, 157.47, 63.30, 61.00, 60.18, 53.41, 26.43, 15.27; MS (EI) m/z 245, 186, 172, 142, 114; Anal. C 48.90, H 6.21, N 5.73 (calcd C 48.98, H 6.16, N 5.71).

Preparation 65.

(Scheme M: M-4 where R₃ is ethyl, Y is CO₂⁻, and the stereochemistry is (2*R*-*trans*))
(2*R*-*trans*)-1,2,4-Azetidinetricarboxylic acid 1-ethyl-2-methyl diester (C₉H₁₃NO₆).

A mixture of M-3 (Scheme M, where R₃ is ethyl, Y is CO₂⁻, and the stereochemistry is
 5 (2*R*-*trans*)) (1.68 g, 6.85 mmol) and LiOH (7.00 mmol) in 1:1 MeOH/H₂O (40 mL) is
 stirred at rt for 45 h, and then is concentrated. The residue is dissolved in half-saturated
 NaHCO₃, and the solution is extracted with Et₂O. The Et₂O solution is discarded. The
 aqueous solution is adjusted with concentrated HCl to pH 4, and is concentrated to a
 yellow solid. This solid is triturated with CHCl₃. The CHCl₃ solution is filtered and
 10 concentrated to give, as a light brown oil, the title compound: TLC (600:400:1
 Hexanes/Acetone/HCO₂H) *R*_f = 0.21-0.45; ¹H NMR (CDCl₃) δ 4.87 (1H), 4.77-4.68 (1H),
 4.64-4.50 (1H), 4.18-3.97 (2H), 3.76 (3H), 2.52-2.33 (2H), 1.26-1.13 (3H); MS (-ESI,
 MeOH solution) *m/z* 230.1. This material is used without purification.

Preparation 66 and Example 140.

15 (Scheme M: M-6 where R₃ is ethyl, R₂ is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is
 CO₂⁻, and the stereochemistry is [2*R*-(2α,4β(*S**))]
[2*R*-(2α,4β(*S))]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-
 oxoethyl]amino]carbonyl]-1,2-azetidinedicarboxylic acid 1-ethyl-2-methyl diester**
 (C₂₆H₂₇Cl₂N₃O₈).

20 To a mixture of Preparation M-4 (1.08 g, 4.67 mmol) and HOBt+H₂O (0.63 g, 4.7 mmol)
 in CH₂Cl₂ (10 mL) at 0 °C is added a solution of EDC (1.04 g, 5.42 mmol) in CH₂Cl₂ (15
 mL). This mixture is stirred at 0 °C for 30 min. It is then treated with M-5 (Scheme M
 where R₃ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*)) (1.88 g,
 4.66 mmol) and *N*-methylmorpholine (0.52 mL, 4.7 mmol). The resulting solution was
 25 stirred at 0 °C for 2 h and at rt for 2 h. The reaction mixture is diluted with 10% KHSO₄
 and extracted with CH₂Cl₂. The combined CH₂Cl₂ extracts are washed with satd NaHCO₃
 and brine, and are combined, dried, filtered and concentrated to give a yellow foam (2.58
 g), that is purified by flash chromatography to give, as a white solid, the title compound:
 mp 97-99 °C; TLC (9:1 CHCl₃/acetone) *R*_f = 0.30; [α]_D²⁵ +79 (*c* 1.02, MeOH); UV
 30 (MeOH) λ_{max} 225 (ε 12200: sh), 251 (17500); ¹³C NMR (CD₃OD) δ 171.50, 171.30,

171.08, 163.67, 156.23, 155.96, 136.92, 136.19, 133.21, 133.03, 131.87, 130.93, 129.64, 129.44, 127.92, 119.97, 61.61, 60.56, 59.82, 59.11, 58.41, 53.45, 51.58, 51.52, 36.57, 36.29, 24.91, 13.43 (22 lines expected; 28 lines observed); MS (FAB) m/z 580.1260 (calcd for $[M+H]^+$, 580.1253); Anal. C 51.95, H 4.56, N 6.89, Cl 15.18 (calcd for 0.61% H₂O: C 53.48, H 4.73, N 7.20, Cl 12.14).

Preparation 67 and Example 141.

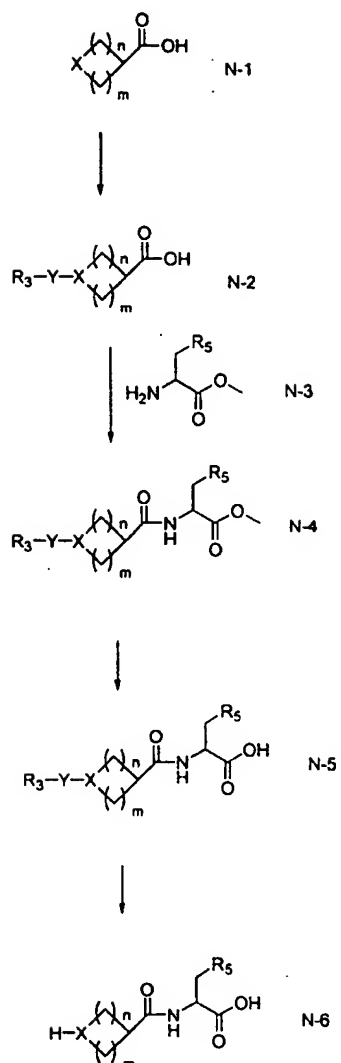
(Scheme M: M-7 where R₃ is ethyl, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is CO₂-, and the stereochemistry is [2*R*-(2 α ,4 β (S*))])

[2*R*-(2 α ,4 β (S*))]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]-amino]-carbonyl]-1,2-azetidinedicarboxylic acid 1-ethyl ester (C₂₄H₂₃Cl₂N₃O₈).

To a suspension of M-6 (Example 140, Scheme M where R₃ is ethyl, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is CO₂-, and the stereochemistry is [2*R*-(2 α ,4 β (S*))]) (0.201 g, 0.347 mmol) in MeOH (5 mL) is added H₂O (4.3 mL) and 1.00 M LiOH (0.70 mL). The reaction mixture is stirred at rt for 23 h. It is concentrated *in vacuo*. The aqueous concentrate is diluted with H₂O, and the solution is adjusted to approximately pH 12 with 1N NaOH. It is extracted with Et₂O, and the Et₂O extract is discarded. The aqueous solution is adjusted to approximately pH 3 with 1N HCl. It is extracted repeatedly with Et₂O. The combined Et₂O extracts were dried, filtered and concentrated to give, as a white solid, the title compound: TLC (500:500:1 MeOH/CH₂Cl₂/HCO₂H) R_f = 0.20; ¹H NMR (CD₃OD) δ 7.59 (2H), 7.50-7.37 (3H), 7.23 (2H), 4.78-4.64 (2H), 4.61-4.51 (1H), 4.22-3.89 (2H), 3.30-3.185 (1H), 3.07-2.93 (1H), 2.42-2.23 (2H), 1.21 and 1.11 (3H total); MS (FAB) m/z 552.0946 (calcd for $[M+H]^+$ 552.0940); Anal. C 50.48, H 4.61, N 6.64, Cl 12.95 (calcd for 1.60% H₂O: C 51.35, H 4.31, N 7.49, Cl 12.63).

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Scheme N.



5

Scheme N teaches a general method for the preparation of *N*-acyl azetidinecarboxylic acid Examples N-5 where *n* equals 0, 1 or 2, *m* equals 0, 1 or 2, (*m* + *n*) equals 2, X is nitrogen and *R*₃, *R*₅ and Y are defined as in Scheme B, and azetidine-carboxylic acid Examples N-6 where *n* equals 0, 1 or 2, *m* equals 0, 1 or 2, (*m* + *n*) equals 2, X is nitrogen and *R*₅ is

10 defined as in Scheme B. Thus acylation of amino acid N-1 gives *N*-acylacid N-2, that is

coupled with reagent N-3 (as exemplified by the use of reagents A-3 and A-4 of Scheme A, and B-4 and B-5 of Scheme B) to provide Examples N-4. Ester hydrolysis of N-4 provides Examples N-5. *N*-deacylation of Examples N-5 provides Examples N-6.

Preparation 68.

5 (Scheme N: N-2 where *n* is 2, *m* is 0, X is N, Y is -CO₂-, R₃ is (1,1-dimethyl)ethyl, and the stereochemistry is (*S*)).

(*S*)-1,2-Azetidinedicarboxylic acid 1-(1,1-dimethylethyl) ester (C₉H₁₅NO₄).

To a mixture of (*S*)-(-)-2-azetidinecarboxylic acid (110 mg, 1.1 mmol), Boc₂O (290 mg, 1.30 mmol), and DMAP (0.017 g, 0.14 mmol) in 4:1 DMF/H₂O (10 mL) is added Et₃N
 10 (0.30 mL, 2.2 mmol). The reaction mixture is stirred at rt for 68 h, and then is concentrated. The concentrate is diluted with EtOAc, and the EtOAc solution is washed with cold 10% KHSO₄. The combined organic extracts are dried, filtered and concentrated to give the title compound as a colorless oil: TLC (750:250:1 Hexanes/acetone/HCO₂H) R_f 0.26; ¹H NMR (CD₃OD) δ 4.97 (1H), 4.57 (1H), 3.98 (1H), 3.87 (1H), 2.57 (H), 2.13
 15 (1H), 1.42 (9H); MS (-ESI) *m/z* 200.3.

Preparation 69 and Example 142.

(Scheme N: N-4 where *n* is 2, *m* is 0, X is N, Y is -CO₂-, R₃ is (1,1-dimethyl)ethyl, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*)).

[2*S*-(*R*^{*},*R*^{*})]-2-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-azetidinecarboxylic acid 1-(1,1-dimethyl)ethyl ester
 20 (C₂₆H₂₉Cl₂N₃O₆).

To a mixture of acid N-2 (Scheme N, where *n* is 2, *m* is 0, X is N, Y is -CO₂-, R₃ is (1,1-dimethyl)ethyl, and the stereochemistry is (*S*)) (1.04 g; 5.17 mmol) and HOBT·H₂O (0.71 g, 5.3 mmol) in CH₂Cl₂ (10 mL) at 0 °C is added a mixture of EDC·HCl (1.00 g, 5.22
 25 mmol) in CH₂Cl₂ (20 mL). The reaction mixture is stirred at 0 °C for 30 min, and then N-3 (Scheme N where R₃ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*)) (2.10 g, 5.20 mmol) and *N*-methylmorpholine (0.60 mL, 5.46 mmol) are added. The reaction mixture is stirred at 0°C for 30 min and at rt for 3 h. The reaction mixture is partitioned between 10% KHSO₄ and CH₂Cl₂. The aqueous phase is extracted twice more
 30 with CH₂Cl₂. The combined organic extracts are washed with saturated NaHCO₃ and brine,

and then are dried, filtered and concentrated to a yellow oil (2.54 g) that is purified by silica flash chromatography to give the title compound as a white foam: mp 106-108 °C; TLC (1:1 Hexanes/EtOAc); R_f = 0.21; $[\alpha]_D^{25}$ -38 (c 1.01, MeOH); ^{13}C NMR (CD_3OD) δ 172.37, 171.58, 163.64, 156.68, 136.98, 136.22, 133.11, 131.87, 130.90, 129.49, 127.91, 120.00, 80.42, 61.87, 53.31, 51.45, 36.40, 27.16, 20.08; MS (EI) m/z 551, 549, 478, 476, 451, 449, 396, 394, 351, 349, 280, 278, 175, 173; Anal. C 56.33, H 5.48, N 7.23, Cl 12.43 (calcd for 0.52% H_2O : C 56.44, H 5.34, N 7.59, Cl 12.82).

Preparation 70 and Example 143.

(Scheme N: N-5 where n is 2, m is 0, X is N, Y is $-\text{CO}_2^-$, R_3 is (1,1-dimethyl)ethyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is [2*S*-(R^* , R^*)])
[2*S*-(R^* , R^*)]-2-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]-carbonyl]-1-azetidinecarboxylic acid 1-(1,1-dimethylethyl) ester ($\text{C}_{25}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_6$).
 A solution of N-4 (Scheme N, where n is 2, m is 0, X is N, Y is $-\text{CO}_2^-$, R_3 is (1,1-dimethyl)ethyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*)) (507 mg, 0.92 mmol) and LiOH (1.25 mmol) in 1:1 MeOH/ H_2O (10 mL) is stirred at rt for 18 h. The reaction mixture is diluted with cold 10% KHSO_4 and extracted with CH_2Cl_2 . The organic extracts are dried, filtered and concentrated to a white foam (498 mg), that is purified by silica flash chromatography to give the title compound: TLC (750:250:1 hexanes/acetone/ HCO_2H) R_f = 0.12; $[\alpha]_D^{25}$ -27 (c 0.94, CHCl_3); ^{13}C NMR (CD_3OD) δ 210.05, 172.70, 172.27, 163.63, 156.75, 136.89, 136.22, 133.36, 131.88, 130.90, 129.56, 127.91, 119.95, 80.47, 61.93, 53.18, 36.47, 27.15, 20.09; MS (-ESI) m/z 533.8; MS (EI) m/z 435, 419, 417, 401, 399, 373, 371, 280, 278, 175, 173, 147, 145; Anal. C 55.23, H 5.25, N 7.42, Cl 12.87 (calcd for 1.10% H_2O : C 55.36, H 5.14, N 7.75, Cl 13.07).

Preparation 71 and Example 144.

(Scheme N: N-6 where n is 2, m is 0, X is N, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (2*S*), L)
N-[[[(2*S*)-2-Azetidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine trifluoroacetic acid salt ($\text{C}_{20}\text{H}_{19}\text{Cl}_2\text{N}_3\text{O}_6 \cdot \text{C}_2\text{HF}_3\text{O}_2$).
 A solution of N-5 (Scheme N, where n is 2, m is 0, X is N, Y is $-\text{CO}_2^-$, R_3 is (1,1-dimethyl)ethyl, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is [2*S*-

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(*R*,R**)) (900 mg; 1.7 mmol) in 1:1 TFA/CH₂Cl₂ (5 mL) is stirred at rt for 1.5 h, and is concentrated. The residue is thrice diluted with CHCl₃ and re-concentrated. This residue is dissolved in MeOH and concentrated to a white foam, that is dissolved in 1:1 MeOH/H₂O and then concentrated to remove most of the MeOH. The solution is frozen and lyophilized to give the product as a white powder: [α]_D²⁵ -6 (*c* 0.72, MeOH); ¹³C NMR (CD₃OD) δ 172.62, 167.54, 163.78, 136.82, 136.13, 133.43, 131.84, 130.97, 129.38, 127.93, 120.15, 58.38, 54.03, 43.73, 36.27, 23.33; MS (+ESI) *m/z* 436.0; MS (FAB) *m/z* 438, 436; Anal. C 46.77, H 3.75, N 7.24, Cl 12.44 (calcd for a 1:1 TFA salt with 1.68% H₂O: C 47.21, H 3.79, N 7.51, Cl 12.67).

Example 145.

(Scheme N: N-4 where *n* is 1, *m* is 1, X is N, R₃ is (1,1-dimethyl)ethyl, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is CO₂⁻ and the stereochemistry is (*S*))

[*S*]-3-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]-carbonyl]-1-azetidinecarboxylic acid 1-(1,1-dimethyl)ethyl ester (C₂₆H₂₉Cl₂N₃O₆).

Example 145 was prepared as described in Scheme N from 3-azetidinecarboxylic acid.

Physical properties as follows: TLC (1:1 EtOAc/hexanes) *R_f* = 0.22; [α]_D²⁵ +18 (*c* 0.92, MeOH); ¹³C NMR (CD₃OD) δ 172.92, 171.91, 163.68, 156.55, 136.87, 136.20, 133.27, 131.89, 130.91, 129.36, 127.90, 120.04, 79.80, 53.85, 51.45, 36.43, 32.26, 27.24; MS (EI) *m/z* 469, 467, 451, 359, 351, 349, 280, 278, 175, 173, 57; Anal. C 56.82, H 5.39, N 7.52, Cl 12.81 (calcd for 0.06% H₂O: C 56.70, H 5.31, N 7.63, Cl 12.87).

Example 146.

(Scheme N: N-5 where *n* is 1, *m* is 1, X is N, R₃ is (1,1-dimethyl)ethyl, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl, Y is CO₂⁻ and the stereochemistry is (*S*))

(*S*)-3-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]]-1-azetidinecarboxylic acid 1-(1,1-dimethylethyl) ester (C₂₅H₂₇Cl₂N₃O₆).

Example 146 was prepared from example 145 by the procedure described in preparation

70. Physical properties as follows: TLC (600:400:1 Hexanes/Acetone/HCO₂H) *R_f* = 0.17; [α]_D²⁵ +33 (*c* 0.92, MeOH); ¹³C NMR (CD₃OD) δ 173.05, 172.89, 163.69, 156.56, 136.78,

136.19, 133.59, 131.89, 130.91, 129.40, 127.90, 120.00, 79.78, 53.67, 51.72, 36.47, 32.31,

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27.24: MS (-ESI) m/z 533.9; MS (FAB) m/z 538, 536, 438, 436, 337, 335, 280, 278, 175, 173, 57; Anal. C 55.03, H 5.21, N 7.52, Cl 12.81 (calcd for 1.22% H₂O: C 55.30, H 5.15, N 7.74, Cl 13.06).

Example 147.

- 5 (Scheme N: N-6 where n is 1, m is 1, X is NH, R₃ is 4-[(2,6-dichlorobenzoyl)amino]-phenyl, and the stereochemistry is (L))

N-[[3-Azetidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine, trifluoroacetic acid salt (C₂₀H₁₉Cl₂N₃O₆ · C₂HF₃O₂).

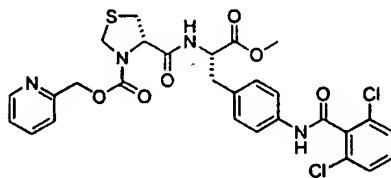
Example 147 was prepared from Example 146 by the procedure described in preparation

- 10 71. Physical properties as follows: $[\alpha]^{25}_D$ +32 (c 0.87, MeOH); ¹³C NMR (CD₃OD) δ 173.32, 170.57, 163.80, 136.70, 136.13, 133.76, 131.84, 130.97, 129.45, 127.93, 120.17, 54.03, 36.59, 35.32; MS (FAB) m/z 438, 436, 391, 331, 175, 173, 101, 55; Anal. C 46.85, H 4.07, N 7.33, Cl 12.39 (calcd for a 1:1 TFA salt with 3.34% H₂O: C 46.41, H 3.92, N 7.38, Cl 12.45).

15

Example 148.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(2-pyridinylmethyl) ester
(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to proton, R₃ is 2-pyridinylmethyl, Y is CO₂-, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and
20 stereochemistry is (S, S)).



- Example 148 was prepared as described in Scheme A from D-cysteine using 2-pyridinemethanol to form the requisite carbamate. Physical data as follows: ¹H NMR (300 MHz, CDCl₃) δ 8.37 (1 H), 7.70 (1 H), 7.51 (2 H), 7.27 (6 H), 7.08 (2 H), 6.92 (1 H), 5.24 (2 H), 4.77 (3 H), 4.40 (1 H), 3.74 (3 H), 3.37 (1 H), 3.15 (3 H); ¹³C NMR (75 MHz,
- 25

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CDCl₃) δ 171.4, 162.5, 155.2, 149.0, 137.1, 136.5, 136.0, 132.4, 130.7, 129.9, 128.0, 123.1, 120.6, 63.0, 53.4, 52.5, 36.8; MS (ESI+) for C₂₈H₂₆Cl₂N₄O₆S *m/z* 616.8 (M+H)⁺; HRMS (EI) calcd for C₂₈H₂₆Cl₂N₄O₆S 616.0950, found 616.0946. Anal. Calcd for C₂₈H₂₆Cl₂N₄O₆S: C, 54.46; H, 4.24; N, 9.07. Found: C, 54.61; H, 4.32; N, 8.97.

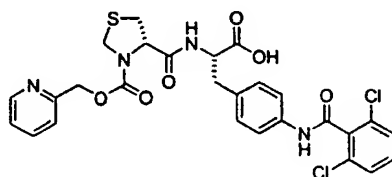
5

Example 149.

[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)-
amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(2-
pyridinylmethyl) ester

10

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to proton, R₃ is 2-pyridinylmethyl, Y is CO₂-, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



15

Example 149 was prepared from example 148 by the procedure described in preparation 6.

Physical data as follows: IR (mull) 1713, 1666, 1605, 1576, 1561, 1539, 1515, 1442, 1431, 1413, 1351, 1325, 1271, 1194, 766 cm⁻¹; ¹H NMR (300 MHz, CD₃OD) δ 8.80 (1 H), 8.56 (1 H), 7.99 (2 H), 7.58 (2 H), 7.45 (3 H), 7.25 (2 H), 5.43 (2 H), 4.60 (3 H), 3.30 (3 H), 2.93 (2 H); ¹³C NMR (75 MHz, CD₃OD) δ 172.7, 171.5, 163.7, 152.8, 151.3, 146.6, 141.9, 136.8, 136.1, 133.6, 131.8, 130.9, 129.6, 127.9, 126.1, 125.4, 120.1, 63.2, 62.8, 59.4, 53.5, 36.4, 35.1; MS (ESI+) for C₂₇H₂₄Cl₂N₄O₆S *m/z* 602.9 (M+H)⁺; MS (FAB) *m/z* (rel. intensity) 603 (MH⁺, 59), 605 (43), 603 (59), 154 (51), 139 (99), 137 (46), 136 (47), 123 (67), 105 (58), 103 (61), 93 (38); HRMS (FAB) calcd for C₂₇H₂₄Cl₂N₄O₆S + H₁ 603.0872, found 603.0876; Anal. Calcd for C₂₇H₂₄Cl₂N₄O₆S • 0.3 H₂O: C, 53.26; H, 4.07; N, 9.20. Found: C, 52.97; H, 4.23; N, 9.04.

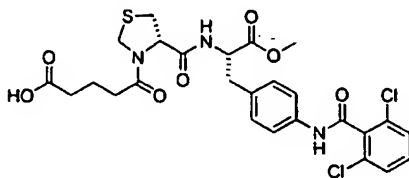
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Example 150.

[S-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]- δ -oxo-3-thiazolidinepentanoic acid

(Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to proton. R₃ is (CH₂)₃CO₂H,

5 Y is CO-, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)).



Example 150 was prepared as described in Scheme A from D-cysteine using glutaric
 10 anhydride to form the requisite amide. Physical data as follows: IR (mull) 3077, 3053, 3040, 1738, 1728, 1696, 1682, 1641, 1557, 1437, 1430, 1414, 1307, 1232, 1209 cm⁻¹; ¹H NMR (300 MHz, CDCl₃) δ 7.56 (2 H), 7.30 (3 H), 7.08 (2 H), 4.96 (1 H), 4.62 (3 H), 3.72 (3 H), 3.28 (4 H), 2.37 (4 H), 1.90 (2 H); ¹³C NMR (75 MHz, CDCl₃) δ 179.5, 176.0, 175.6, 173.6, 167.2, 140.6, 140.0, 136.1, 134.5, 133.7, 133.4, 131.8, 124.4, 66.7, 65.9,
 15 57.2, 56.3, 40.3, 39.3, 37.3, 36.7, 36.2, 33.5, 23.6; MS (FAB) *m/z* (rel. intensity) 596 (MH⁺, 90), 598 (63), 597 (40), 596 (90), 341 (25), 263 (25), 230 (32), 225 (31), 193 (31), 141 (99), 88 (36); HRMS (FAB) calcd for C₂₆H₂₇Cl₂N₃O₇S + H₁, 596.1025, found 596.1036. Anal. Calcd for C₂₆H₂₇Cl₂N₃O₇S • 0.3 H₂O: C, 51.88; H, 4.62; N, 6.98. Found: C, 51.69; H, 4.69; N, 6.59.

20

Example 151.

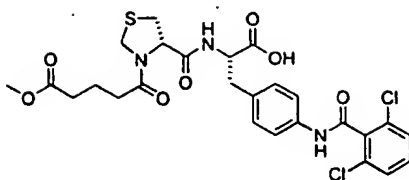
[S-(R*,R*)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]- δ -oxo-3-thiazolidinepentanoic acid methyl ester

(Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to proton. R₃ is

(CH₂)₃CO₂CH₃, Y is CO-, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and

25 stereochemistry is (S, S)).

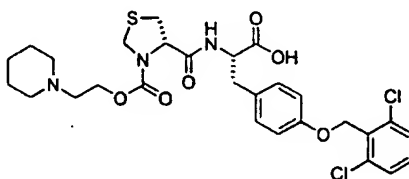
-185-



Example 151 was prepared as described in Scheme A from D-cysteine using methyl glutaryl chloride to form the requisite amide. Physical data as follows: IR (mull) 3287, 3196, 1724, 1662, 1607, 1562, 1540, 1516, 1431, 1414, 1326, 1268, 1217, 1195, 799 cm^{-1} ;
 5 MS (FAB) m/z (rel. intensity) 596 (MH^+ , 72), 598 (52), 596 (72), 229 (37), 193 (37), 167 (34), 133 (44), 129 (69), 121 (48), 103 (83), 89 (99); ^1H NMR (300 MHz, CD_3OD) δ 7.56 (2 H), 7.43 (3 H), 7.23 (2 H), 4.53 (3 H), 3.63 (3 H), 2.96 (4 H), 2.45 (3 H), 2.24 (2 H), 1.93 (2 H); ^{13}C NMR (75 MHz, CDCl_3) δ 174.0, 172.1, 169.9, 162.9, 162.8, 136.1, 132.2,
 10 130.6, 130.0, 127.9, 120.3, 62.9, 54.5, 51.6, 49.8, 36.8, 33.5, 33.0, 32.7, 29.6, 19.6; HRMS (FAB) calcd for $\text{C}_{26}\text{H}_{27}\text{Cl}_2\text{N}_3\text{O}_7\text{S} + \text{H}$, 596.1025, found 596.1047. MS (FAB) m/z (rel. intensity) 596 (MH^+ , 72), 598 (52), 596 (72), 229 (37), 193 (37), 167 (34), 133 (44), 129 (69), 121 (48), 103 (83), 89 (99).

Example 152.

15 $[S-(R^*,R^*)]-4-[[[1\text{-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]-carbonyl]-3\text{-thiazolidinecarboxylic acid } 3\text{-[2-(1-piperidiny)ethyl] ester}$
 (Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to proton, R_3 is 2-(1-piperidiny)ethyl, Y is CO_2^- , R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).
 20



Example 152 was prepared from example 5 by the procedure described in preparation 6. Physical data as follows: IR (mull) 3254, 2654, 1711, 1565, 1547, 1512, 1438, 1344,

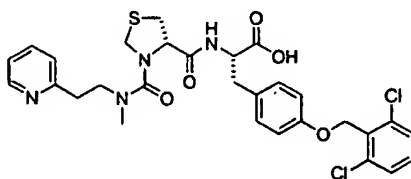
-186-

1300, 1240, 1196, 1179, 1119, 1014, 767 cm^{-1} ; ^1H NMR (300 MHz, CD_3OD) δ 7.45 (2 H), 7.36 (1 H), 7.19 (2 H), 6.97 (2 H), 5.26 (2 H), 4.50 (5 H), 3.60 (11 H), 1.83 (6 H); MS (ESI+) for $\text{C}_{38}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 610.0 ($\text{M}+\text{H}$) $^+$; Anal. Calcd for $\text{C}_{38}\text{H}_{33}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 1.5 \text{H}_2\text{O} \cdot \text{HCl}$: C, 49.90; H, 5.53; N, 6.24; Cl, 15.78. Found: C, 49.86; H, 5.43; N, 6.29; Cl, 15.65.

5 % Water (KF): 3.99.

Example 153.

[*S*-(*R* * ,*R* *)]-4-[[[1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]-carbonyl]-*N*-methyl-*N*-[2-(2-pyridinyl)ethyl]-3-thiazolidinecarboxamide
(Scheme A, A-8: where $\text{R}_{\text{A-1}}$ and $\text{R}_{\text{A-2}}$ are the same and equal to proton, R_3 is 2-(2-pyridyl)-ethyl, Y is $\text{CON}(\text{CH}_3)-$, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*, *S*)).

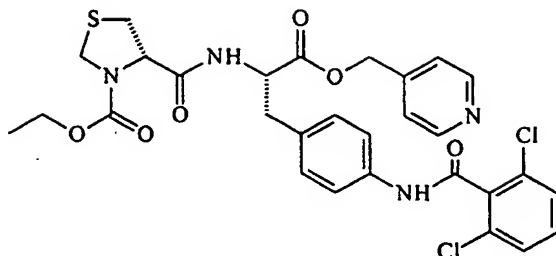


15 Example 153 was prepared as described in Scheme A using 2-(2-methylaminoethyl)-pyridine to form the requisite urea and hydrolysis according to the procedure described in preparation 6. Physical data as follows: mp 80°C (softens), 125°C; IR (mull) 1661, 1611, 1585, 1565, 1511, 1489, 1439, 1394, 1300, 1240, 1196, 1179, 1017, 779, 768 cm^{-1} ; ^1H NMR (300 MHz, CD_3OD) δ 8.44 (1 H), 7.75 (1 H), 7.35 (5 H), 7.12 (2 H), 6.93 (2 H),
20 5.22 (2 H), 4.83 (1 H), 4.65 (1 H), 4.32 (2 H), 3.77 (1 H), 3.45 (1 H), 3.20 (1 H), 3.00 (5 H), 2.84 (3 H); ^{13}C NMR (75 MHz, CD_3OD) δ 173.0, 170.8, 162.4, 158.4, 157.9, 147.9, 138.0, 136.6, 132.2, 130.6, 130.1, 129.3, 128.3, 124.2, 122.0, 114.5, 64.8, 64.7, 53.3, 52.5, 49.7, 35.7, 34.8, 32.8; MS (ESI+) for $\text{C}_{29}\text{H}_{30}\text{Cl}_2\text{N}_4\text{O}_5\text{S}$ m/z 617.0 ($\text{M}+\text{H}$) $^+$; Anal. Calcd for $\text{C}_{29}\text{H}_{30}\text{Cl}_2\text{N}_4\text{O}_5\text{S}$: C, 56.40; H, 4.90; N, 9.07. Found: C, 56.31; H, 5.07; N, 8.98.

25 Preparation 72 and Example 154.

[*S*-(*R* * ,*R* *)]-4-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-[(4-pyridinyl)-methoxy]-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester

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To a solution of Example 12 (Scheme A, A-8: where R_{A-1} and R_{A-2} are the same and equal to H, R_3 is ethyl, Y is CO_2 , R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (S, S)) (400 mg, 0.74 mmol) in dimethylformamide (4 mL) was added tetramethylguanidine (204 μL , 1.63 mmol) followed by 4-picolyl chloride (138 mg, 0.81 mmol). The solution was heated to 65 $^\circ\text{C}$ for 3 h and volatiles removed *in vacuo*.

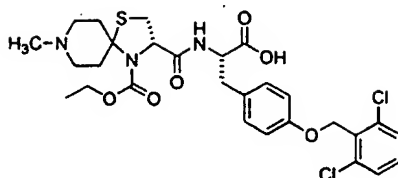
Purification of the residue by flash chromatography using methylene chloride/methanol (2%) as eluant afforded the title compound (320 mg) as an amorphous solid: IR (mull)

3275, 1748, 1677, 1608, 1561, 1539, 1515, 1431, 1415, 1344, 1325, 1271, 1222, 1194, 799 cm^{-1} ; ^1H NMR (300 MHz, CDCl_3) δ 8.87 (1 H), 8.48 (2 H), 7.50 (2 H), 7.25 (3 H), 7.10 (2 H), 6.93 (2 H), 5.08 (2 H), 4.79 (2 H), 4.61 (1 H), 4.28 (1 H), 4.13 (2 H), 3.16 (4 H), 1.21 (3 H); ^{13}C NMR (75 MHz, CDCl_3) δ 174.6, 170.7, 170.6, 162.6, 149.2, 144.5, 136.9, 136.0, 132.3, 131.7, 130.7, 129.8, 128.0, 122.5, 120.3, 65.0, 63.0, 62.7, 53.3, 37.3, 20.9, 14.5; MS (ESI+) for $\text{C}_{29}\text{H}_{28}\text{Cl}_2\text{N}_4\text{O}_6\text{S}$ m/z 630.8 (M+H) $^+$; Anal. Calcd for $\text{C}_{29}\text{H}_{28}\text{Cl}_2\text{N}_4\text{O}_6\text{S}$: C, 55.15; H, 4.47; N, 8.87. Found: C, 54.85; H, 4.58; N, 8.74. Anal. Calcd for $\text{C}_{29}\text{H}_{28}\text{Cl}_2\text{N}_4\text{O}_6\text{S}$: C, 55.15; H, 4.47; N, 8.87; Cl, 11.23; S, 5.08. Found: C, 54.85; H, 4.58; N, 8.74.

Preparation 73 and Example 155.

[S-(R^*, R^*)]-3-[[[1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]-carbonyl]-8-methyl-1-thia-4,8-diazaspiro[4.5]decane-4-carboxylic acid 4-ethyl ester (Scheme B, B-7: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} together form a cyclic ring of 6 atoms of the formula $-\text{CH}_2\text{CH}_2\text{N}(\text{CH}_3)\text{CH}_2\text{CH}_2-$, Y is CO_2 , R_3 is ethyl, R_5 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).

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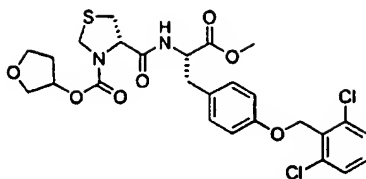


- t*-Butyl ester B-6 (Scheme B where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} together form a cyclic ring of 6 atoms of the formula -CH₂CH₂N(CH₃)CH₂CH₂-, Y is
- 5 CO₂, R₃ is ethyl, R_{B-5} is O-*t*-butyl, R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*, *S*), prepared according to Scheme B from 1-methyl-4-piperidone and D-cysteine) (681 mg, 1.02 mmol) was dissolved in a solution of HCl in dioxane (4 M, 28 mL) at ambient temperature. After 18 h, volatiles were removed *in vacuo* to afford a residue (650 mg) which was lyophilized from water. Further purification of a portion of
- 10 this product (200 mg) was effected by chromatography on a Biotage Flash 40™ system using a 40 g KP-C18-HS (35-70 μm) silica gel cartridge using aqueous acetonitrile (40%) as eluant to afford the title compound (94 mg) as an amorphous powder: IR (mull) 1696, 1611, 1585, 1565, 1511, 1439, 1404, 1335, 1303, 1271, 1239, 1196, 1178, 1017, 769 cm⁻¹; ¹H NMR (DMSO-d₆, 300 MHz) δ 8.25 (1 H), 7.54 (2 H), 7.43 (1 H), 7.13 (2 H), 6.94 (2 H), 5.16 (2 H), 4.84 (1 H), 4.41 (1 H), 4.00 (2 H), 3.07 (4 H), 2.73 (4 H), 2.53 (3 H), 1.96 (1 H), 1.69 (1 H), 1.11 (3 H); ¹³C NMR (CDCl₃, 75 MHz) δ 173.9, 169.2, 157.7, 153.2, 136.9, 132.1, 130.7, 130.5, 129.3, 128.5, 114.7, 73.2, 66.7, 65.2, 62.8, 54.0, 43.5, 36.5, 14.6; MS (ESI+) for C₂₈H₃₃Cl₂N₃O₆S *m/z* 610.0 (M+H)⁺; MS (ESI-) for C₂₈H₃₃Cl₂N₃O₆S *m/z* 607.9 (M-H)⁻; HRMS (FAB) calcd for C₂₈H₃₃Cl₂N₃O₆S+H₁ 610.1545, found 610.1561.
- 15 Anal. Calcd for C₂₈H₃₃Cl₂N₃O₆S • 0.6 HCl • H₂O: C, 51.70; H, 5.52; N, 6.46; Cl, 14.17. Found: C, 51.28; H, 5.49; N, 6.50; Cl, 14.57. % Water (KF): 2.72.
- 20

Example 156.

- [*S*-(R*,R*)]-4-[[[1-[[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(3-tetrahydrofuranyl) ester
- 25 (Scheme A, A-7: where R_{A-1} and R_{A-2} are the same and equal to proton, R₃ is 3-tetrahydrofuranyl, Y is CO₂-, R₅ is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is (*S*, *S*)).

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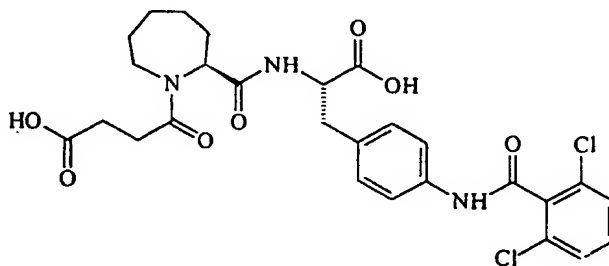


Example 156 was prepared as described in Scheme A from D-cysteine using 3-hydroxytetrahydrofuran to form the requisite carbamate. Physical properties as follows:

mp 125-126.5°C. IR (mull) 3311, 1750, 1744, 1708, 1661, 1549, 1515, 1439, 1408, 1307, 1243, 1227, 1212, 1173, 1019 cm^{-1} ; ^1H NMR (300 MHz, CD_3OD) δ 7.38 (3 H), 7.14 (2 H), 7.96 (2 H), 5.20 (1 H), 5.25 (2 H), 4.61 (4 H), 3.79 (4 H), 3.74 (3 H), 3.19 (2 H), 2.84 (2 H), 2.29 (2 H); ^{13}C NMR (75 MHz, CDCl_3 , spectra complicated via the presence of diastereomers) δ 171.7, 171.6, 158.1, 137.0, 132.1, 130.5, 130.4, 130.3, 128.5, 128.2, 115.1, 77.2, 77.2, 73.2, 67.0, 65.2, 63.1, 63.0, 53.3, 52.5, 52.4, 36.9, 32.9; MS (ESI+) for $\text{C}_{26}\text{H}_{28}\text{Cl}_2\text{N}_2\text{O}_7\text{S}$ m/z 582.8 ($\text{M}+\text{H}$) $^+$; Anal. Calcd for $\text{C}_{26}\text{H}_{28}\text{Cl}_2\text{N}_2\text{O}_7\text{S}$: C, 53.52; H, 4.84; N, 4.80. Found: C, 53.34; H, 4.87; N, 4.86.

Example 157.

[S-(R*,R*)]-2-[[[1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-hexahydro- γ -oxo-1H-azepine-1-butanoic acid



Example 157 was prepared as described for the preparation of Example 167. Physical properties as follows: IR (mull) 1781, 1709, 1651, 1625, 1612, 1550, 1537, 1515, 1444, 1431, 1418, 1398, 1331, 1193, 798 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 1.45 (8 H), 2.30 (4 H), 2.90 (3 H), 3.80 (1 H), 4.50 (2 H), 7.16 (2 H), 7.50 (5 H), 7.94 (1 H), 10.62

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(1 H), 7.71 (1 H): MS (FAB) m/z (rel. intensity) 578 (M+H, 43), 581 (9), 580 (29), 579 (19), 578 (43), 577 (10), 227 (11), 226 (99), 198 (18), 173 (9), 98 (46).

Preparation 74 and Example 158.

(Scheme N, N-6: where n is 2, m is 0, X is N, R_1 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and stereochemistry is 2*S*-(R^* , R^*))
 5 **[2*S*-(R^* , R^*)]-2-[[[1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]azetidine** ($C_{21}H_{21}Cl_2N_3O_4$).

A solution of the product of example 142 (Scheme N: N-4 where n is 2, m is 0, X is N, Y is $-CO_2^-$, R_1 is (1,1-dimethyl)ethyl, R_2 is 4-[(2,6-dichlorobenzoyl)amino]phenyl and
 10 stereochemistry is (*S*)) (512 mg; 0.93 mmol) in 1:1 TFA/ CH_2Cl_2 (10 mL) is stirred at rt for 1 h. The reaction mixture is concentrated under reduced pressure. The residue is taken up in a mixture of CH_2Cl_2 and satd aqueous $NaHCO_3$. The aqueous phase is extracted twice additionally with CH_2Cl_2 . The combined CH_2Cl_2 portions are dried, filtered and concentrated to give a yellow oil (440 mg), that is purified by silica flash chromatography
 15 (95:5 CH_2Cl_2 /MeOH) to afford the title compound (324 mg) as a white foam: mp 113-115°C; TLC (95:5 CH_2Cl_2 /MeOH) R_f = 0.10; $[\alpha]_D^{25}$ -34 (c 0.96, MeOH); UV (MeOH) λ_{max} 224 (sh, ϵ 12100), 251 (17700), 284 (sh, 2880); IR (mineral oil mull) 3260, 1744, 1664, 1606, 1561, 1537, 1515, 1431, 1414, 1323, 1270, 1223, 1195, 799, 782 cm^{-1} ; 1H NMR (CD_3OD) δ 7.61 (2H), 7.50-7.38 (3H), 7.25 (2H), 4.86 (1H), 4.75 (1H), 4.21 (1H),
 20 3.73 (3H), 3.61 (2H), 3.39-3.28 (1H), 3.23 (1H), 3.05 (1H), 2.63-2.50 (1H), 2.21-2.08 (1H); MS (+ESI) m/z 450.0; MS (EI) m/z 451, 449, 396, 394, 351, 349, 278, 211, 175, 173, 96, 70, 56; Anal. C 55.68, H 4.79, N 8.96, Cl 15.43 (calcd for +1.08% H_2O : C 55.41, H 4.77, N 9.23, Cl 15.58).

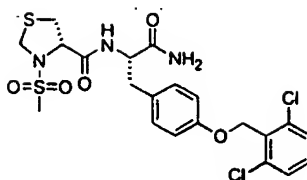
Example 159.

25 ***O*-[(2,6-Dichlorophenyl)methyl]-*N*-[[[(4*S*)-3-(methylsulfonyl)-4-thiazolidinyl]carbonyl]-*L*-tyrosinamide**

(Scheme C, C-10: where R_{C-1} , R_{C-2} , R_{C-3} and R_{C-4} are the same and equal to proton, R_1 is methyl, Y is SO_2^- , R_2 is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (*S*, *S*)).

30

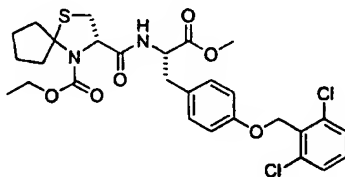
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Example 159 was prepared as described in Scheme C using methanesulfonyl chloride to form the requisite sulfonamide. Physical properties as follows: mp 228-230°C; ¹H NMR (300 MHz, DMSO-d₆) δ 8.03 (1 H), 7.54 (2 H), 7.44 (2 H), 7.14 (3 H), 6.92 (2 H), 5.16 (2 H), 4.69 (2 H), 4.41 (1 H), 4.31 (1 H), 3.21 (1 H), 3.01 (3 H), 2.87 (3 H); ¹³C NMR (75 MHz, DMSO-d₆) δ 172.9, 169.1, 157.5, 136.5, 132.3, 132.0, 130.8, 130.6, 129.2, 114.6, 65.3, 64.4, 54.3, 52.0, 37.3, 35.0; MS (ESI-) for C₂₁H₂₃Cl₂N₃O₅S₂ m/z 530.2 (M-H)⁻; Anal. Calcd for C₂₁H₂₃Cl₂N₃O₅S₂: C, 47.37; H, 4.35; N, 7.89. Found: C, 47.43; H, 4.46; N, 7.81.

Example 160.

[S-(R*,R*)]-3-[[[1-[[4-[(2,6-Dichlorophenyl)methoxy]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-thia-4-azaspiro[4.4]nonane-4-carboxylic acid 4-ethyl ester (Scheme B, B-6: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} together form a carbocyclic ring of 5 atoms, R_{B-5} is OCH₃, Y is CO₂, R₃ is ethyl, R₅ is 4-[(2,6-dichlorophenyl)methoxy]phenyl and stereochemistry is (S, S)).



Example 160 was prepared as described in Scheme B. Physical properties as follows: IR (mull) 1746, 1705, 1681, 1510, 1439, 1399, 1336, 1301, 1276, 1241, 1203, 1178, 1110, 1017, 769 cm⁻¹; ¹H NMR (CDCl₃) δ 7.36 (2 H), 7.25 (1 H), 7.03 (2 H), 6.93 (2 H), 6.64 (1 H), 5.22 (2 H), 4.85 (2 H), 4.15 (2 H), 3.73 (3 H), 3.15 (4 H), 2.69 (1 H), 2.48 (1 H), 1.76 (6 H), 1.23 (3 H); ¹³C NMR (CDCl₃) δ 171.5, 170.7, 158.0, 152.5, 137.0, 132.0, 130.5, 130.3, 128.5, 128.2, 115.0, 66.4, 65.2, 61.9, 53.1, 52.4, 37.1, 32.3, 25.1, 24.6, 14.5; MS (ESI+) for C₂₈H₃₂Cl₂N₂O₆S m/z 594.9 (M+H)⁺. MS (ESI-) for C₂₈H₃₂Cl₂N₂O₆S m/z

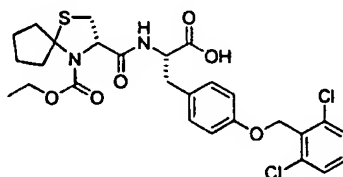
592.8 (M-H)⁺; Anal. Calcd for C₂₈H₃₂Cl₂N₂O₆S • 0.10 H₂O: C, 56.30; H, 5.43; N, 4.69.

Found: C, 56.20; H, 5.24; N, 4.69. % Water (KF): 0.31.

Example 161.

[S-(R*,R*)]-3-[[[1-Carboxy-2-[4-[(2,6-
5 dichlorophenyl)methoxy]phenyl]ethyl]amino]carbonyl]-1-thia-4-azaspiro[4.4]nonane-4-
carboxylic acid 4-ethyl ester

(Scheme B, B-7: where R_{B-1} and R_{B-2} are the same and equal to H, R_{B-3} and R_{B-4} together form a carbocyclic ring of 5 atoms, Y is CO₂, R₃ is ethyl, R₅ is 4-[(2,6-dichlorophenyl)-methoxy]phenyl and stereochemistry is (S, S)).

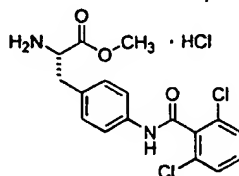


10

Example 161 was prepared from example 160 by the procedure described in preparation 6.

Physical properties as follows: IR (mull) 1737, 1708, 1675, 1612, 1511, 1439, 1402, 1338, 1301, 1241, 1197, 1179, 1115, 1018, 769 cm⁻¹; ¹H NMR (DMSO-d₆) δ 8.07 (1 H), 7.54 (2 H), 7.45 (1 H), 7.12 (2 H), 6.94 (2 H), 5.15 (2 H), 4.63 (1 H), 4.33 (1 H), 3.91
15 (2 H), 3.05 (2 H), 2.79 (1 H), 2.60 (1 H), 1.60 (6 H), 1.07 (3 H); ¹³C NMR (DMSO-d₆) δ 172.7, 169.4, 157.1, 155.7, 135.9, 131.6, 131.4, 130.3, 129.6, 128.7, 114.1, 83.9, 64.7, 60.5, 53.1, 38.0, 36.2, 31.8, 24.2, 24.1, 14.1; MS (ESI⁺) for C₂₇H₃₀Cl₂N₂O₆S m/z 580.8 (M+H)⁺; MS (ESI⁻) for C₂₇H₃₀Cl₂N₂O₆S m/z 578.8 (M-H)⁻; HRMS (EI) calcd for C₂₇H₃₀Cl₂N₂O₆S 580.1202, found 580.1172; Anal. Calcd for C₂₇H₃₀Cl₂N₂O₆S • 0.19 H₂O:
20 C, 55.44; H, 5.24; N, 4.79. Found: C, 55.24; H, 5.32; N, 4.79. % Water (KF): 0.59.

Preparation 75.



The aminoester product of preparation 75 is useful as a synthetic intermediate (for example, reagent A-4 of Scheme A).

To a cold (0-5°C) solution of anhydrous methanolic HCl was added 100 g of L-4-nitrophenylalanine (Advanced ChemTech) portionwise over 15 min. The mechanically stirred mixture was heated to gentle reflux for 48 h. The mixture was allowed to cool and then filtered through a sintered glass filter funnel, washing the collected solids with hot
5 MeOH until only insoluble residues remained. The filtrate was concentrated *in vacuo* to afford the methyl ester (120 g) as waxy off white solid which was used without further purification.

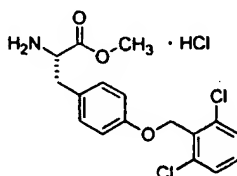
To a suspension of methyl ester described above (87 g, 0.33 mole) in CH₂Cl₂ (1500 mL) at ambient temperature was added di-*t*-butyldicarbonate (109 g, 0.50 mole) followed by the
10 dropwise addition of Et₃N (51 mL, 0.37 mole). After 15 min additional Et₃N (40 mL, 0.29 mol) was added to maintain a slightly basic mixture (*ca.* pH 8). The reaction mixture was stirred 18 h and additional CH₂Cl₂ (1400 mL) and Et₃N (15 mL, 0.11 mol) were added. After an additional 2 h the reaction mixture was quenched by the slow addition of MeOH (100 mL), stirred for 1 h and then partitioned between CH₂Cl₂ and cold 10% aqueous
15 KHSO₄. The organic layer was washed with saturated NaHCO₃ and brine, dried (Na₂SO₄), filtered and concentrated *in vacuo*. Flash chromatography of the residue using hexane and a gradient of a 1:1 mixture of EtOAc/CH₂Cl₂ (25-33%) afforded the Boc-methyl ester (69 g) as a white solid. Physical properties as follows: ¹H NMR (300 MHz; CDCl₃) δ 8.16 (2H), 7.31 (2H), 5.04 (1H), 4.63 (1H), 3.73 (3H), 3.18 (2H), 1.41 (9H); MS (ES+) for
20 C₁₅H₂₀N₂O₆ *m/z* 325.2 (M+H)⁺.

Palladium on carbon (10% w/w, 1.25 g) was added to a Parr hydrogenation flask under a N₂ atmosphere and carefully wetted with 100 mL of MeOH/THF (1:1). A solution of the Boc-methyl ester described above (25 g, 77 mmol) in 400 mL of MeOH/THF (1:1) was added and the mixture shaken on a hydrogenation apparatus under a hydrogen atmosphere
25 (20 psi) for 1 h at ambient temperature. The reaction mixture was filtered through a pad of Celite and the solids washed several times with MeOH. The combined filtrates were concentrated *in vacuo* to afford the 4-aminophenylalanyl derivative (22.7 g) which was used without further purification. Physical properties as follows: ¹H NMR (300 MHz, CDCl₃) δ 6.89 (2H), 6.61 (2H), 4.96 (1H), 4.50 (1H), 3.69 (3H), 2.95 (2H), 1.41 (9H); MS
30 (ES+) for C₁₃H₂₂N₂O₄ *m/z* 295.2 (M+H)⁺.

A cold (0-5°C) solution of 2,6-dichlorobenzoyl chloride (11.1 mL, 77.5 mmol) in 125 mL of THF was treated dropwise with a solution of the 4-aminophenylalanyl derivative described above (22.7 g, 77.1 mmol) and Et₃N (16 mL, 115 mmol) in 125 mL of THF. The reaction mixture was allowed to warm to temperature and stir an additional 18 h. The mixture was diluted with EtOAc (2 L) and then washed with 1N HCl, H₂O, 1N NaOH and brine. The organic extract was dried (Na₂SO₄), filtered, and concentrated *in vacuo* to give the crude product as a pale yellow solid. This material was recrystallized from acetone/hexanes (*ca.* 1:1) to afford the amide (30.8 g) as a crystalline solid. Physical properties as follows: mp 192.2-193.1°C; IR (mull) 3305, 1747, 1736, 1690, 1665, 1609, 1548, 1512, 1433, 1414, 1325, 1277, 1219, 1171, 781 cm⁻¹; ¹H NMR (300 MHz; CDCl₃) δ 7.57 (2H), 7.34 (4H), 7.14 (2H), 4.98 (1H), 4.60 (1H), 3.74 (3H), 3.11 (2H), 1.42 (9H); MS (ES+) for C₂₂H₂₄Cl₂N₂O₃, *m/z* 467.0 (M+H)⁺.

To 650 mL of anhydrous 4M HCl in dioxane at ambient temperature was added the amide described above (30.6 g, 65.5 mmol) portionwise and the resulting mixture was stirred until all the solids had dissolved (*ca.* 1 h). Volatiles were removed *in vacuo* to give a light yellow solid which was partitioned between water (500 mL) and ether (1 L). The water layer was separated and concentrated *in vacuo* to approximately 200 mL. The aqueous solution was then frozen and lyophilized to afford the aminoester product (25.6 g) as a light yellow solid. Physical properties as follows: [α]_D²⁵ = +5 (*c* 1, MeOH); IR (mull) 3244, 3186, 3112, 1747, 1660, 1604, 1562, 1539, 1516, 1431, 1416, 1327, 1273, 1243, 799 cm⁻¹; ¹H NMR (300 MHz; CD₃OD) δ 7.69 (2H), 7.45 (3H), 7.29 (2H), 4.34 (1H), 3.83 (3H), 3.21 (2H); ¹³C NMR (300 MHz; CD₃OD) δ 169.0, 163.9, 137.8, 136.08, 131.8, 131.0, 130.3, 129.7, 127.9, 120.6, 53.8, 52.3, 35.4; MS (ES+) for C₁₇H₁₆Cl₂N₂O₃, *m/z* 367.1 (M+H)⁺.

Preparation 76.



The aminoester product of preparation 75 is useful as a synthetic intermediate (for example, reagent A-4 of Scheme A).

To a cold (0-5°C) solution of anhydrous methanolic HCl (200 mL) was added 25 g of *N*- α -*t*-Boc-*O*-2,6-dichlorobenzyl-L-tyrosine (Sigma) portionwise over 15 min. After 30 minutes at 0-5°C, the mixture was heated to 50°C for 2 h. The solution was cooled to room temperature and the volatiles removed *in vacuo*. The solid was suspended in ethyl ether and collected by filtration to afford the title compound (21.4 g) which was used without further purification. Physical properties as follows: $[\alpha]_D^{25} = +16$ (c 1.00, ethanol); ¹H NMR (300 MHz, CD₃OD) δ 7.44 (2 H), 7.35 (1 H), 7.21 (2 H), 7.02 (2 H), 5.28 (2 H), 4.29 (1 H), 3.81 (3 H), 3.18 (2 H); MS (ESI+) for C₁₇H₁₇Cl₂NO₃, *m/z* 354.1 (M+H)⁺; Anal. Calcd for C₁₇H₁₇Cl₂NO₃•HCl: C, 52.26; H, 4.64; N, 3.59. Found: C, 52.17; H, 4.74; N, 3.61.

Example 162

2-[[[(1S)-1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-1-piperidinecarboxylic acid 1-[(1,1-dimethylethyl) ester



Example 162: HCl gas was bubbled through a solution of *N*-(tert-butoxycarbonyl)-4-(2,6-dichlorobenzoylamino)-L-phenylalanine (2.51g, 5.53 mmol) in MeOH (20 mL) for 10 minutes. The solution was stirred for additional 2 hours at room temperature. The solvent was removed *in vacuo* and the excess HCl was removed by the addition of Et₂O (3 x 15 mL) and evaporation under reduced pressure. The resultant gum was dissolved in THF (10 mL) and *N*-tert-butoxycarbonyl-pipecolinic acid (1.28 gm, 5.59 mmol), BOP-reagent

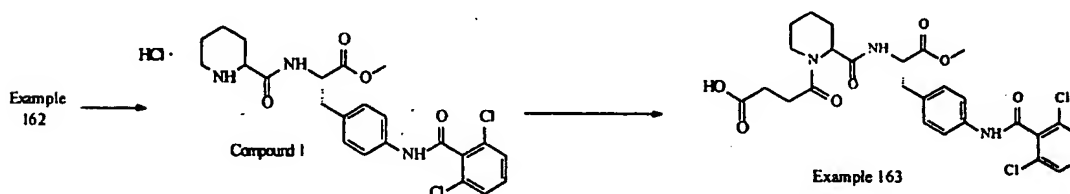
-196-

(2.69 gm, 6.09 mmol) and DIEA (2.9 mL, 16.6 mmol) were added and the reaction mixture was stirred overnight. EtOAc (25 mL) was added and the mixture was extracted with 1N HCl (20 mL). The organic phase was washed with saturated LiCl (20 mL) then saturated NaHCO₃ (30 mL). The organic layer was dried over Na₂SO₄, filtered and
 5 evaporated. Chromatography of the residue (silica gel, Hexanes -> 50% EtOAc/Hexanes gradient elution) provided Example 162 as a solid (1.45 gm, 45 %): ESMS (m/z) 578, 580 (MH⁺).

Example 163

2-[[[(1S)-1-[[4-[(2,6-Dichlorobenzoyl)amino]phenyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-γ-oxo-1-piperidinebutanoic acid

10



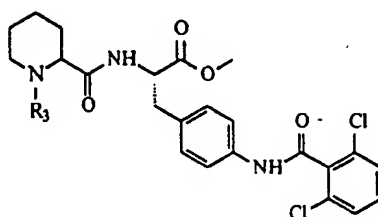
HCl gas was bubbled through a MeOH (20 mL) solution of Example 162 (1.27 gm, 2.20 mmol) for 10 min. Stirring was continued overnight at room temperature. The solvent was removed *in vacuo* and the excess HCl was removed by washing with Et₂O (3
 15 x 10 mL) on a vacuum filter. The HCl salt was completely dried under high vacuum to provide Compound 1 (1.09 gm, 97 %) as a solid: ESMS (m/z) 478, 480 (MH⁺).

Example 163: Compound 1 (147 mg, 0.285 mmol) was dissolved in DMF (5 mL) containing DIEA (150 μL, 0.88 mmol). To this solution was added succinic anhydride (59
 20 mg, 0.59 mmol) and the mixture was stirred at 50°C for 5 hr under dry nitrogen. The solvent was evaporated and the residue was purified by column chromatography (silica gel, Hexanes -> EtOAc gradient elution) to provide Example 163 as a solid (164 mg): ESMS (m/z) 578, 580 (MH⁺).

Examples 164-166

The following mono methyl esters were prepared in a similar manner as Example 163

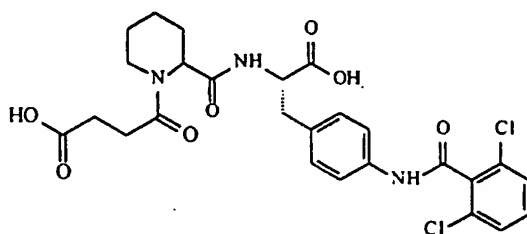
5



ex#	R ₃	MS (m/z)
164		646 (MH ⁺)
165		620 (MH ⁺)
166		574 ([M-H] ⁻)

Example 167

2-[[[(1S)-1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-γ-oxo-1-piperidinebutanoic acid



Example 167

Example 163 (154 mg, 0.266 mmol) was treated with LiOH (26 mg, 1.07 mmol) in H₂O (5 mL) for 3 hours. The product was then precipitated by the addition of 3 N HCl.

15

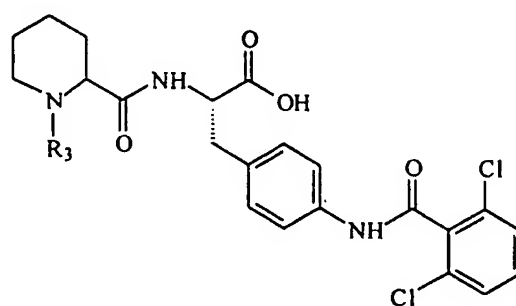
The product was collected by vacuum filtration and washed with cold H₂O (2 x 3 mL).

Drying under high vacuum provided Example 167 as a solid (109 mg): ESMS (m/z) 562 ([M-H]⁻).

5

Example 168-170

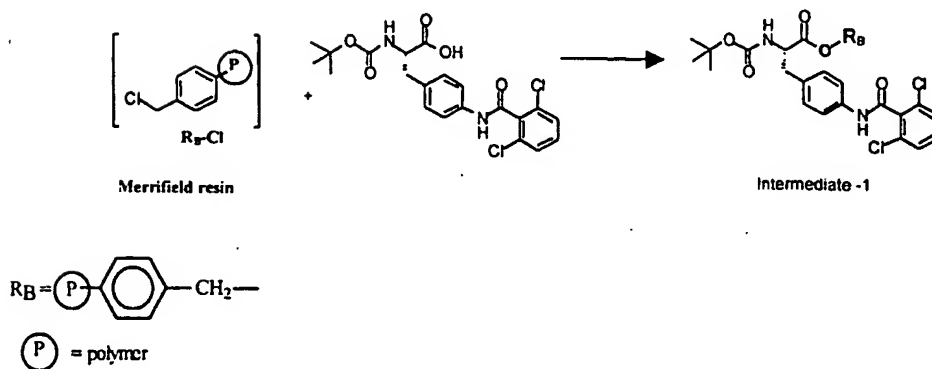
The following compounds were prepared in a similar manner as 167.



ex#	R ₃	MS (m/z) ([M-H] ⁻)
168		630
169		604
170		560

10

Preparation 77

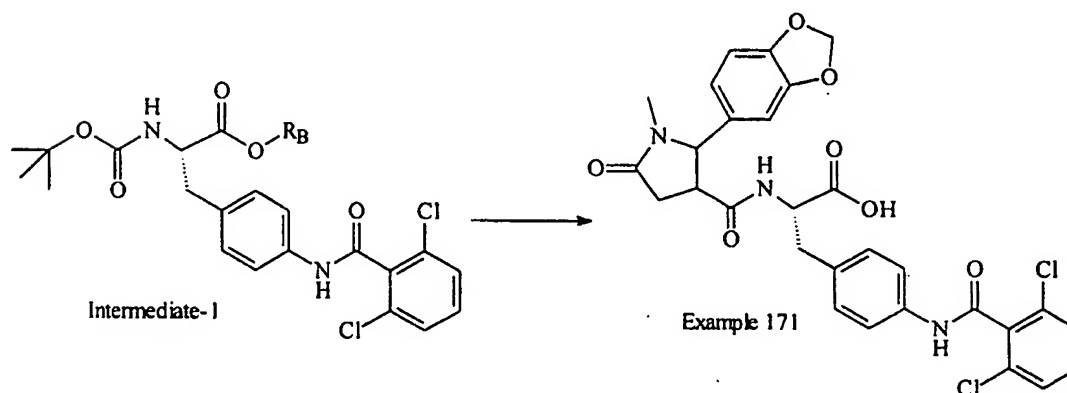


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Intermediate-1: Attachment of N-tert-butoxycarbonyl-[4-(2,6-dichlorobenzoylamino)]-L-phenylalanine to Merrifield resin was done using Horiki's method (Horiki et al., *Chem. Lett.* 1978(2) 165-168). In a 250 mL round bottom flask fitted with a drying tube, Merrifield resin (Biorad, 10.0 g, 13.5 mmol/g) and anhydrous potassium fluoride (Aldrich, 1.57g, 27.0 mmol) were added to a solution of N-tert-butoxycarbonyl-[4-(2,6-dichlorobenzoylamino)]-L-phenylalanine (Bachem California, 6.13 g, 13.5 mmol) in dry DMF (100 mL). The reaction mixture was stirred at 80°C in an oil bath for 24 hr. The cooled resin was then filtered and washed thoroughly with DMF (2 x 250 mL), 50% aqueous DMF (3 x 250 mL), methanol (3 x 250 mL), dichloromethane (3 x 250 mL), and finally methanol (3 x 250 mL). The resin was then dried under reduced pressure to constant weight to give Intermediate-1. Incorporation of N-tert-butoxycarbonyl-[4-(2,6-dichlorobenzoylamino)]-L-phenylalanine onto the resin was estimated to be 0.045 mmol/g from the increase in resin mass.

Example 171

N-[[2-(1,3-Benzodioxol-5-yl)-1-methyl-5-oxo-3-pyrrolidinyl]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine



Example 171: The **Intermediate-1** (0.15 g, 0.1065 mmol/g) was pretreated with CH₂Cl₂ (2x 3 mL). The swollen resin was then deprotected with 50% TFA/CH₂Cl₂ (3mL, 30min). The resin was rinsed in the following order: CH₂Cl₂ (2 x 3 mL), CH₃OH (2 x 3 mL), CH₂Cl₂ (2 x 3 mL). The resin was swollen with DMF (2 x 3 mL). 2-(3,4-

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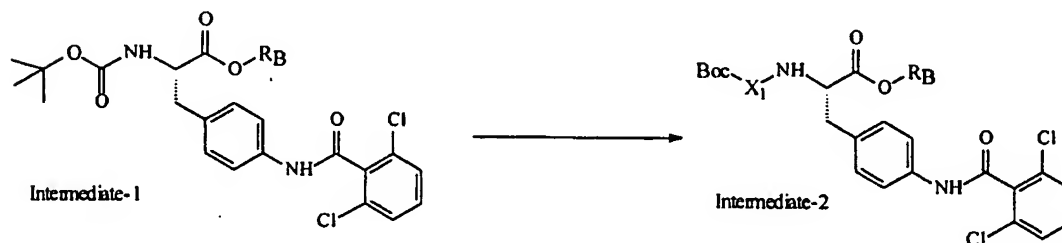
methylenedioxyphenyl)-1-methyl-5-oxo-3-pyrrolidine carboxylic acid (84 mg, 0.32 mmol) in DMF (1.0 mL) was activated with 0.5 M HBTU/HOBT in DMF (0.7 mL) and DIEA (0.139 mL, 0.799 mmol), and then added to the swollen resin. The mixture was vortexed for 2 hr at room temperature. The resin was filtered and washed in the following order:

5 DMF(2 x 3 mL), CH₂Cl₂ (2 x 3 mL), CH₃OH (2 x 3 mL), CH₂Cl₂ (2 x 3 mL), respectively. If a Kaiser test on a small quantity of the resin is positive (blue) then repeat the coupling procedure until a negative result is obtained. The resulting resin was then dried *in vacuo* to constant weight. The resin was placed in the polypropylene column and pretreated with 3 mL of THF. Then to the preswollen resin 1.6 mL of THF, 0.48 mL of

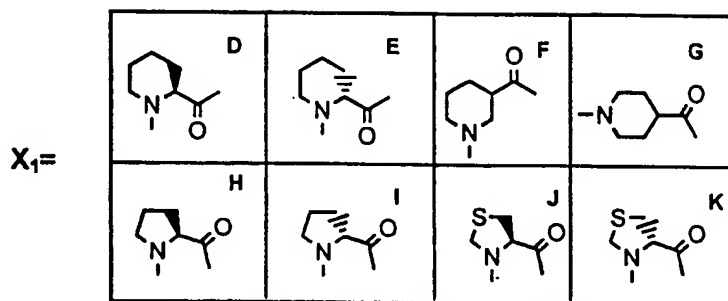
10 CH₃OH, and 0.160 mL of 2N LiOH were added. The mixture was vortexed for 15 min and filtered to a clean and preweighed test tube. The resin was next washed with 2mL of THF/5% CH₃OH (2x) and the combined filtrates were evaporated. The resulting gum was dissolved in 1 mL of water. The solution was then acidified with 1N HCl to pH 2.0. The precipitate was centrifuged, washed with water (5mL, 2x) and dried *in vacuo* to furnish

15 15.4 mg of Example 171 as a solid. ESMS (m/z): 596, ([M-H]⁻).

Preparation 78



-201-

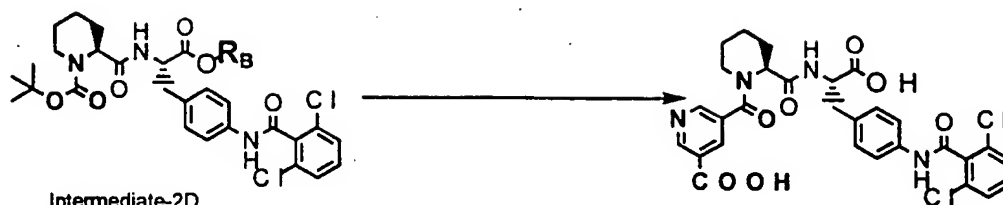


Intermediate-2F: The resin bound N-tert-butoxycarbonyl-[4-(2,6-dichlorobenzoylamino)]-L-phenylalanine (**Intermediate-1**), (250 mg, 0.1125 mmol/g) was placed in a 8.0 mL, polypropylene filter column fitted with a 2-way polypropylene stopcock. The resin was pretreated with CH_2Cl_2 (2 x 3 mL). The swollen resin was then deprotected with 50% TFA/ CH_2Cl_2 (3-4 mL, 30min) with shaking. The resin was rinsed in the following order: CH_2Cl_2 (2 x 3 mL), CH_3OH (2 x 3 mL), CH_2Cl_2 (2 x 3 mL). The resin was swollen with DMF (2 x 3 mL). N-Tert-butoxycarbonyl-nipicotic acid (103 mg, 0.45 mmol) in DMF (1.0 mL) was activated with 0.5 M HBTU/HOBT in DMF (0.910 mL) and DIEA (0.195 mL), then added to the swollen resin. The mixture was vortexed for 2 hr at room temperature. The resin was washed in the following order: DMF (2 x 3 mL), CH_2Cl_2 (2 x 3 mL), CH_3OH (2 x 3 mL), CH_2Cl_2 (2 x 3 mL) and dried (**Intermediate-2F**). If a Kaiser test on a small quantity of the resin is positive (blue) then repeat the coupling procedure until a negative result is obtained. The intermediate resins Intermediate-2D, 2E, 2G, 2H, 2I, 2J & 2K were each produced following this procedure.

Example 172

5-[[[(2S)-2-[[[(1S)-1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-1-piperidinyl]carbonyl]-3-pyridinecarboxylic acid

-202-



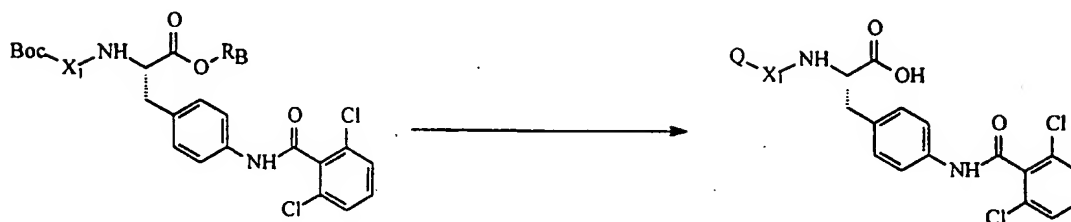
Example 172

Example 172: Resin bound **Intermediate-2D** (2.0 g, 1.3 mmol/g) was pretreated with CH_2Cl_2 (2 x 20 mL). The swollen resin was then deprotected with 50% TFA/ CH_2Cl_2 (20 mL, 30min). The resin was rinsed in the following order: CH_2Cl_2 (2 x 20 mL), CH_3OH (2 x 20 mL), CH_2Cl_2 (2 x 20 mL). The resin was swollen with DMF (2 x 20 mL). 3,5-Pyridine dicarboxylic acid (652 mg, 3.9 mmol) in 20 mL of DMF was activated with 0.5 M HBTU/HOBT in DMF (8.0 mL) and DIEA (1.7 mL, 9.75 mmol), then added to the swollen resin. The mixture was vortexed for 2 hr at room temperature. The resin was filtered and washed in the following order: DMF (2 x 20 mL), CH_2Cl_2 (2 x 20 mL), CH_3OH (2 x 20 mL), CH_2Cl_2 (2 x 20 mL), respectively. If a Kaiser test (Kaiser et al., *Anal. Biochem.* 1970, 34, 594-598) on a small quantity of the resin is positive (blue) then repeat the coupling procedure until a negative result is obtained. The resulting resin was then dried *in vacuo* to constant weight (2.2 g). The resin was treated with 25 mL of liquid HF by stirring for 60 min at 0°C in an HF-reaction apparatus (Peninsula Laboratories Inc., Belmont, CA). The HF was rapidly evaporated off by vacuum aspiration at 0°C. Then 100 mL of dry ethyl ether was added. The resin and the resulting precipitates were filtered off and washed three times with 50 mL of ethyl ether, and dried *in vacuo*. The mixture was then treated with 25 mL of 1N NaOH (4x), and the combined solutions were lyophilized. The crude product was purified by HPLC using a C-18 column and a linear acetonitrile/0.1% HCl gradient. The gradient was run from 60% solvent A (0.1% HCl) to

80% solvent B (80% acetonitrile in 0.1% HCl) in 20 min. Lyophilization furnished 20 mg (2.5%) of Example 172. ESMS (m/z): 612 ($[M-H]^-$).

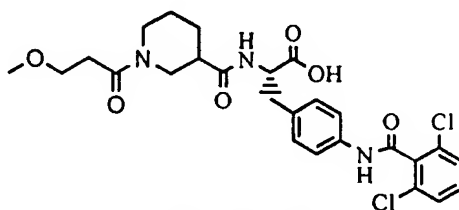
Examples 173-267

5



Method-A (Example 173)

10 4-[(2,6-Dichlorobenzoyl)amino]-*N*-[[1-(3-methoxy-1-oxopropyl)-3-piperidinyl]carbonyl]-
L-phenylalanine



Example 173

15

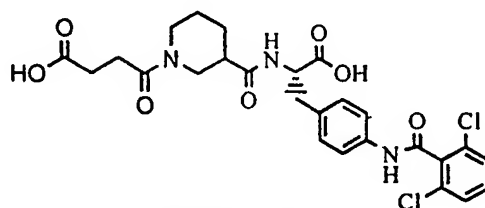
The **Intermediate-2F** (0.25 g, 0.1125 mmol/g) was pretreated with CH_2Cl_2 (2 x 3 mL). The swollen resin was then deprotected with 50% TFA/ CH_2Cl_2 (3mL, 30min). The resin was rinsed in the following order: CH_2Cl_2 (2 x 3 mL), CH_3OH (2 x 3 mL), CH_2Cl_2 (2 x 3 mL). The resin was swollen with DMF (2 x 3 mL). 3-methoxypropionic acid (53 mg, 0.45 mmol) in DMF (1.0 mL) was activated with 0.5 M HBTU/HOBT in DMF (0.910 mL) and DIEA (0.195 mL), then added to the swollen resin. The mixture was vortexed for 2 hr at room temperature. The resin was filtered and washed in the following order: DMF

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(2 x 3 mL), CH₂Cl₂ (2 x 3 mL), CH₃OH (2 x 3 mL), CH₂Cl₂ (2 x 3 mL), respectively. If a Kaiser test on a small quantity of the resin is positive (blue) then repeat the coupling procedure until a negative result is obtained. The resulting resin was then dried *in vacuo* to constant weight. The resin was placed in the polypropylene column and pretreated with THF (3 mL). Then THF (3.5 mL), CH₃OH (1.0 mL) and 2N LiOH (0.175 mL) were added. The mixture was vortexed for 15 min and filtered to a clean and preweighed test tube. The resin was next washed with THF/5% CH₃OH (2mL) and the combined filtrates were evaporated. The resulting gum was dissolved in H₂O (1 mL). The solution was then acidified with 1N HCl to pH 2.0. The precipitate was centrifuged, washed with water (2 x 5 mL) and dried *in vacuo* to furnish 38.3 mg of Example 173 as a solid: ESMS (m/z) 548 ([M-H]⁻).

Method B (Example 174):

3-[[[(1S)-1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-γ-oxo-1-piperidinebutanoic acid



Example 174

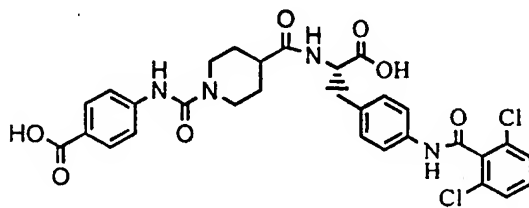
The **Intermediate-2F** (0.25g, 0.1125 mmol/g) was pretreated with CH₂Cl₂ (2 x 3 mL). The swollen resin was then deprotected with 50% TFA/CH₂Cl₂ (3mL, 30min). The resin was rinsed in the following order: CH₂Cl₂ (2 x 3 mL), CH₃OH (2 x 3 mL), CH₂Cl₂ (2 x 3 mL). The resin was then swollen with DMF (3mL). Succinic anhydride (45 mg, 0.45 mmol) dissolved in DMF (4 mL) was added to the swollen resin and stirred at 50°C for 2hr. The resin was filtered and washed in the following order: DMF (2 x 3 mL), CH₂Cl₂ (2 x 3 mL), CH₃OH (2 x 3 mL), CH₂Cl₂ (2 x 3 mL), respectively. If a Kaiser

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test on a small quantity of the resin is positive (blue) then repeat the coupling procedure until a negative result is obtained. The resulting resin was then dried *in vacuo* to constant weight. The resin was placed in the polypropylene column and pretreated with THF (3 mL). Then to the swollen resin THF (3.5 mL), CH₃OH (1.0 mL) and 2N LiOH (0.175 mL) were added. The mixture was vortexed for 15 min and filtered to a clean and preweighed test tube. The resin was next washed with THF/5% CH₃OH (2 x 2 mL) and the combined filtrates were evaporated. The resulting gum was dissolved in H₂O (1 mL). The solution was then acidified with 1N HCl to pH 2.0. The precipitate was centrifuged, washed with H₂O (2 x 5 mL) and dried *in vacuo* to furnish 30.5 mg of Example 174 as a solid: ESMS (m/z) 562 ([M-H]⁻).

Method C (Example 175):

N-[[1-[[[(4-Carboxyphenyl)amino]carbonyl]-4-piperidiny]carbonyl]-4-[(2,6-dichlorobenzoyl)amino]-L-phenylalanine



Example 175

The **Intermediate-2G** (0.25g, 0.1125 mmol/g) was pretreated with CH₂Cl₂ (2 x 3 mL). The swollen resin was then deprotected with 50% TFA/CH₂Cl₂ (3mL, 30min). The resin was rinsed in the following order: CH₂Cl₂ (2 x 3 mL), CH₃OH (2 x 3 mL), CH₂Cl₂ (2 x 3 mL). The resin was then swollen with DMF (3mL). Ethyl 4-isocyanatobenzoate (22 mg, 0.108 mmol) dissolved in DMF (3 mL) and DIEA (47 μL 0.27 mmol) were added to the swollen resin. This reaction mixture was vortexed for 6-8 hr at room temperature. The resin was filtered and washed in the following order: DMF (2 x 3 mL), CH₂Cl₂ (2 x 3 mL), CH₃OH (2 x 3 mL), CH₂Cl₂ (2 x 3 mL), respectively. If a Kaiser test on a small

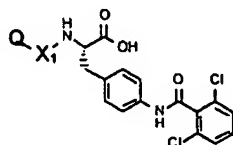
quantity of the resin is positive (blue) then repeat the coupling procedure until a negative result is obtained. The resulting resin was then dried *in vacuo* to constant. The resin was placed in the polypropylene column and pretreated with THF (3 mL). Then to the swollen resin THF (3.5 mL), CH₃OH (1.0 mL) and 2N LiOH (0.175 mL) were added respectively.

- 5 The mixture was vortexed for 15 min and filtered to a clean and preweighed test tube. The resin was next washed with THF/5% CH₃OH (2mL) and the combined filtrates were evaporated. The resulting gum was dissolved in H₂O (1 mL). The solution was then acidified with 1N HCl to pH 2.0. The precipitate was centrifuged, washed with H₂O (2 x 5 mL) and dried *in vacuo* to furnish 18 mg of Example 175 as a solid: ESMS (m/z) 625, 10 ([M-H]⁻).

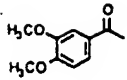





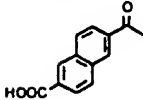
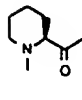
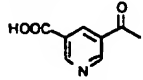





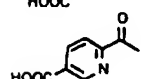

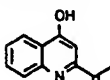

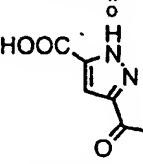
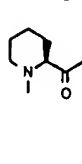
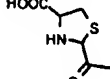

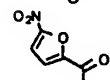





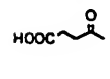

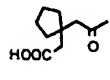

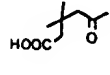

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

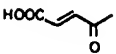

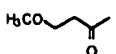





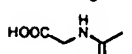

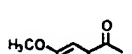





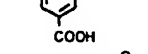

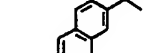

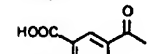

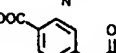

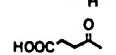

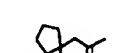

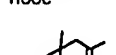

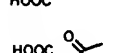

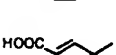

The following compounds were prepared in a similar manner as described above.


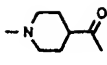

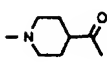
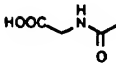
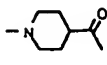
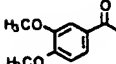
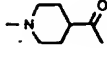

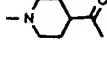

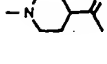
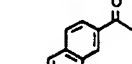
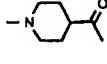
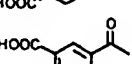
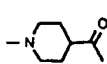
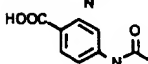
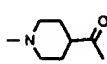


















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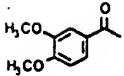







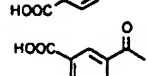

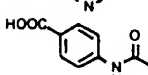





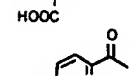

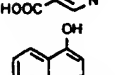

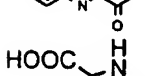







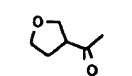

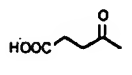

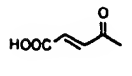



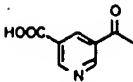

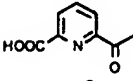





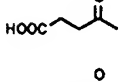

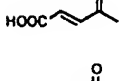

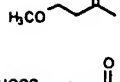

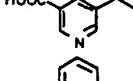

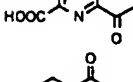





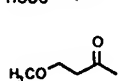

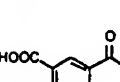

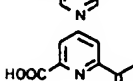

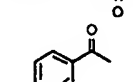





ex#	Method	Q	X ₁	MS (m/z) ([M-H] ⁻)
176	A			560 ([M-H] ⁻)
177	A			548 ([M-H] ⁻)
178	A			616 ([M-H] ⁻)
179	A			560 ([M-H] ⁻)

180	A			626 ([M-H] ⁻)
181	A			610 ([M-H] ⁻)
182	A			610 ([M-H] ⁻)
183	A			660 ([M-H] ⁻)
184	A			611 ([M-H] ⁻)
185	A			567 ([M-H] ⁻)
186	A			611 ([M-H] ⁻)
187	A			611 ([M-H] ⁻)
188	A			633 ([M-H] ⁻)
189	A			600 ([M-H] ⁻)
190	A			621 ([M-H] ⁻)
191	A			603 ([M-H] ⁻)
192	A			560 ([M-H] ⁻)
193	A			560 ([M-H] ⁻)
194	B			562 ([M-H] ⁻)
195	B			630 ([M-H] ⁻)
196	B			604 ([M-H] ⁻)

197	B			560 ([M-H] ⁻)
198	A			560 ([M-H] ⁻)
199	A			548 ([M-H] ⁻)
200	A			616 ([M-H] ⁻)
201	A			560 ([M-H] ⁻)
202	C			563 ([M-H] ⁻)
203	A			626 ([M-H] ⁻)
204	A			610 ([M-H] ⁻)
205	A			610 ([M-H] ⁻)
206	A			660 ([M-H] ⁻)
207	A			611 ([M-H] ⁻)
208	C			625 ([M-H] ⁻)
209	B			562 ([M-H] ⁻)
210	B			630 ([M-H] ⁻)
211	B			604 ([M-H] ⁻)
212	B			560 ([M-H] ⁻)
213	A			560 ([M-H] ⁻)
214	A			548 ([M-H] ⁻)

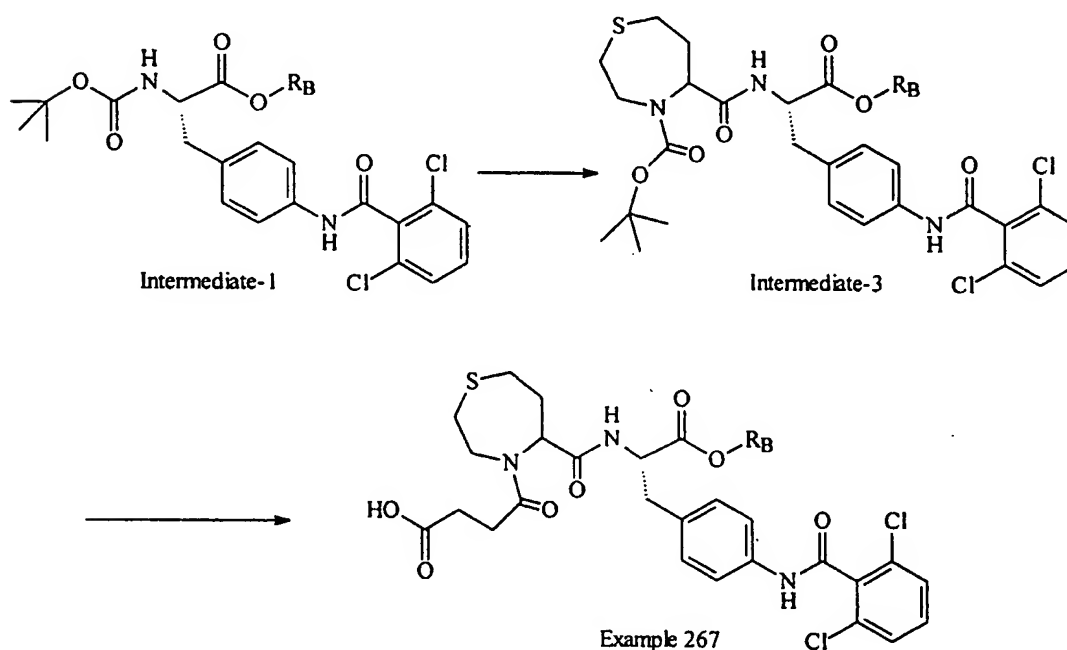
215	A			616 ([M-H] ⁻)
216	A			560 ([M-H] ⁻)
217	C			563 ([M-H] ⁻)
218	A			626 ([M-H] ⁻)
219	A			610 ([M-H] ⁻)
220	A			610 ([M-H] ⁻)
221	A			660 ([M-H] ⁻)
222	A			611 ([M-H] ⁻)
223	C			625 ([M-H] ⁻)
224	B			548 ([M-H] ⁻)
225	B			616 ([M-H] ⁻)
226	B			590 ([M-H] ⁻)
227	B			546 ([M-H] ⁻)
228	A			546 ([M-H] ⁻)
229	A			534 ([M-H] ⁻)
230	A			602 ([M-H] ⁻)
231	A			546 ([M-H] ⁻)
232	C			549 ([M-H] ⁻)

233	A			612 ([M-H] ⁻)
234	A			596 ([M-H] ⁻)
235	A			596 ([M-H] ⁻)
236	A			646 ([M-H] ⁻)
237	A			597 ([M-H] ⁻)
238	C			611 ([M-H] ⁻)
239	A			571 ([M-H] ⁻)
240	A			615 ([M-H] ⁻)
241	A			615 ([M-H] ⁻)
242	A			637 ([M-H] ⁻)
243	A			604 ([M-H] ⁻)
244	A			625 ([M-H] ⁻)
245	A			564 ([M-H] ⁻)
246	A			564 ([M-H] ⁻)
247	A			562 ([M-H] ⁻)
248	A			560 ([M-H] ⁻)
249	A			548 ([M-H] ⁻)

250	A			611 ([M-H] ⁻)
251	A			611 ([M-H] ⁻)
252	A			567 ([M-H] ⁻)
253	A			560 ([M-H] ⁻)
254	A			548 ([M-H] ⁻)
255	A			546 ([M-H] ⁻)
256	A			534 ([M-H] ⁻)
257	A			597 ([M-H] ⁻)
258	A			597 ([M-H] ⁻)
259	A			553 ([M-H] ⁻)
260	A			546 ([M-H] ⁻)
261	A			564 ([M-H] ⁻)
262	A			552 ([M-H] ⁻)
263	A			615 ([M-H] ⁻)
264	A			615 ([M-H] ⁻)
265	A			571 ([M-H] ⁻)
266	A			564 ([M-H] ⁻)

Example 267

5-[[[(1S)-1-Carboxy-2-[4-[(2,6-dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-tetrahydro- γ -oxo-1,4-thiazepine-4(5H)-butanoic acid

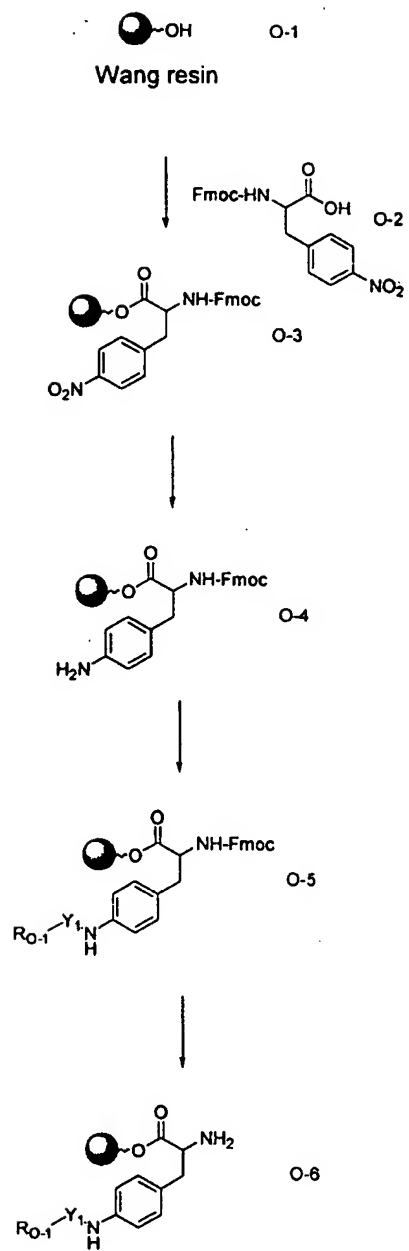


- 5 The **Intermediate-1** (0.3g, 0.195mmol/g) was pretreated with CH_2Cl_2 (2x 3 mL). The swollen resin was then deprotected with 50% TFA/ CH_2Cl_2 (3mL, 30 min). The resin was rinsed in the following order: CH_2Cl_2 (2x 3 mL), CH_3OH (2x 3mL), CH_2Cl_2 (2x 3mL). The resin was swollen with DMF (2x 3mL). N-Tert-butoxycarbonyl-1,4-thiazoline-5-carboxylic acid (204 mg, 0.78 mmol) in DMF (2.0 mL) was activated with 0.5 M
- 10 HBTU/HOBT in DMF (1.6 mL) and DIEA (0.340 mL, 1.95 mmol), then added to the swollen resin. The mixture was vortexed for 2 hr at room temperature. The resin was filtered and washed in the following order: DMF (2x 3 mL), CH_2Cl_2 (2x 3mL), CH_3OH (2x 3mL), CH_2Cl_2 (2x 3mL), respectively. If a Kaiser test on a small quantity of the resin is positive (blue) then repeat the coupling procedure until a negative result is obtained. The
- 15 resin (intermediate-3, 0.2 g, 0.13mmol/g) was pretreated with CH_2Cl_2 (2x 3 mL). The swollen resin was then deprotected with 50% TFA/ CH_2Cl_2 (3mL, 30 min). The resin was rinsed in the following order: CH_2Cl_2 (2x 3mL), CH_3OH (2x 3mL), CH_2Cl_2 (2x 3mL). The resin was then swollen with DMF (3mL). Succinic anhydride (78 mg, 0.78 mmol)

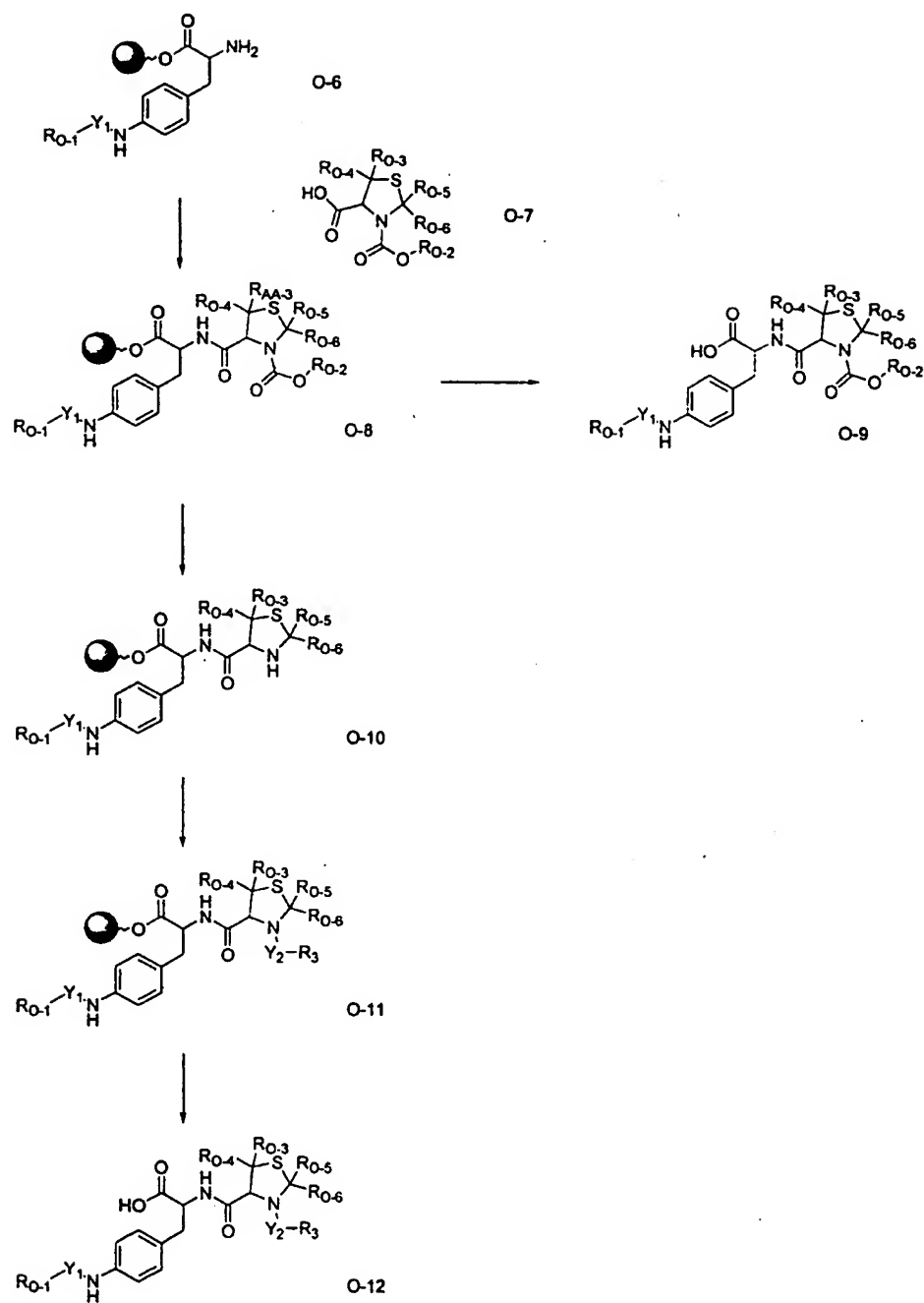
dissolved in DMF (4mL) was added to the swollen resin and stirred at 50°C for 2hr. Then the resin was filtered and washed in the following order: DMF (2x 3mL), CH₂Cl₂ (2x 3mL), CH₃OH (2x 3mL), CH₂Cl₂ (2x 3mL) respectively. If a Kaiser test on a small quantity of the resin is positive (blue) then repeat the coupling procedure until a negative
5 result is obtained. The resulting resin was then dried in vacuo to constant weight. The resin was placed in the polypropylene column and pretreated with THF (3 mL). Then THF (3.9 mL), CH₃OH (1.2 mL) and 2N LiOH (0.195 mL) were added to the swollen resin. The mixture was vortexed for 15 min and filtered to a clean and preweighed test tube. The resin was next washed with THF/5% CH₃OH (2x 2mL) and the combined filtrates were
10 evaporated. The resulting gum was dissolved in H₂O (1 mL). The solution was then acidified with 1N HCl to pH 2.0. The precipitate was centrifuged, washed with H₂O (2 x 5mL) and dried in vacuo to furnish 80 mg of Example 267 as a solid: ESMS (m/z) 594([M-H]⁻).

-214-

Scheme O



Scheme O (continued)



Where: R_{O-1} is defined as R_{12} ; R_{O-2} is defined as C_{1-6} alkyl or C_{7-17} arylalkyl; R_{O-3} , R_{O-4} , and R_{O-5} , are defined independently as R_1 . R_{O-6} is defined as R_2 . Y_1 and Y_2 are defined independently as Y .

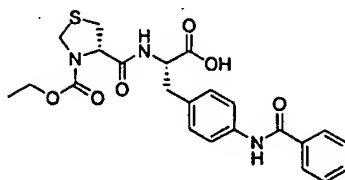
5 Scheme O describes a method for the preparation of examples of the formula **O-9** and **O-12**. Commercially available Wang resin (**O-1**) is acylated with commercially available *N*- α -Fmoc-Phe(NO₂)-OH (**O-2**) under standard conditions to afford the resin of formula **O-3**. Reduction of the aromatic nitro group (Meyer *et al.*, *Mol. Diversity* 1995, 1, 13-20) affords the resin bound aniline (**O-4**) which may be reacted with a variety of
10 electrophilic reagents to afford resin bound amides (**O-5** where Y_1 is C(=O)), ureas (**O-5** where Y_1 is C(=O)NH), sulfonamides (**O-5** where Y_1 is SO₂), and carbamates (**O-5** where Y_1 is C(=O)O). Removal of the Fmoc group under standard conditions provides amine of general structure **O-6** which is acylated using standard solid-phase peptide synthesis conditions (Atherton, E.; Sheppard R.C. *Solid Phase Peptide Synthesis: A Practical*
15 *Approach*; IRL Press at Oxford University Press: Oxford, 1989) with a commercially available or readily prepared thiazolidine-4-carboxylic acid of general formula **O-7** to afford the resin bound intermediate **O-8**. Mild cleavage under standard conditions (Atherton, E.; Sheppard R.C. *Solid Phase Peptide Synthesis: A Practical Approach*; IRL Press at Oxford University Press: Oxford, 1989) affords the acid of general structure **O-9**.
20 In those cases where R_{O-2} is a 9-fluorenylmethyl group, standard Fmoc group removal affords the amine of general structure **O-10**, which may be reacted with a variety of electrophilic reagents as described in Scheme A to afford resin bound amides, ureas, sulfonamides and carbamates of general structure **O-11**. Mild cleavage under standard conditions affords the acid of general structure **O-12**.

25

Preparation 79 and Example 268.

(4*S*)-4-[[[(1*S*)-1-Carboxy-2-[4-[(benzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester

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To a mixture of Wang resin (1% DVB, Advanced Chemtech, 2.75 g, 2.20 mmol based on manufacture's loading of 0.8 mmol/g resin) in DMF (12 mL) was added *N*- α -Fmoc-Phe(NO₂)-OH, **O-2** (Advanced Chemtech, 1.90 g, 4.40 mmol) at room temperature.

5 After mixing for 10 min (by passing a slow stream of nitrogen through the mixture) pyridine (587 μ L, 7.26 mmol) and 2,6-dichlorobenzoyl chloride (630 μ L, 4.40 mmol) were added. The mixture was agitated overnight via nitrogen bubbling, filtered, washed with DMF, methylene chloride and methanol and dried *in vacuo*. In order to cap any unreacted hydroxymethyl groups, the resin was suspended in dichloroethane (5 mL) and to this
10 mixture was added benzoyl chloride (0.75 mL) and pyridine (0.75 mL). The mixture was agitated for 2 h, filtered, washed with DMF, methylene chloride, methanol, methylene chloride and methanol, and dried *in vacuo* to afford the resin **O-3** (3.30 g). IR (diamond anvil) 1733, 1606 (resin), 1520 (resin), 1494 (resin), 1452 (resin), 1347, 1247, 1174, 1029 cm⁻¹.

15 To the prewashed (2 X 20 mL DMF) resin **O-3** (1.0 g, ca. 0.6 mmol based on an adjusted loading of 0.6 mmol/g) was added SnCl₂ • 2 H₂O (6 mL of a 2M solution in DMF, 12 mmol). The viscous suspension was agitated for 4 hours by nitrogen bubbling, filtered and washed with DMF (2 X 20 mL). The resin was resuspended with SnCl₂ • 2 H₂O (6 mL of a 2M solution in DMF, 12 mmol), agitated overnight via nitrogen bubbling, filtered,
20 washed extensively with DMF, water, 2-propanol, methylene chloride and methanol and dried *in vacuo* to afford the resin **O-4**. Examination of the FTIR spectra of a small sample of resin **O-4** failed to exhibit an absorption at 1347 cm⁻¹.

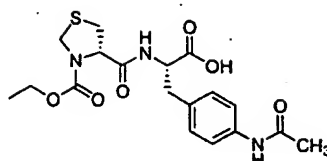
To a mixture of prewashed (2 X 20 mL CH₂Cl₂) resin **O-4** (0.30 g, ca. 0.18 mmol based on an adjusted loading of 0.6 mmol/g) in 1,2-dichloroethane (3 mL) was added
25 benzoyl chloride (174 μ L, 1.50 mmol) and DIEA (313 μ L, 1.80 mmol). The mixture was agitated overnight via nitrogen bubbling, filtered, washed with methylene chloride, DMF, methanol, and methylene chloride and dried *in vacuo* to afford resin **O-5**. To a mixture of

resin O-5 in methylene chloride (5 mL) was added a solution of piperidine in DMF (30%, 5 mL). A slow stream of nitrogen was bubbled through the mixture to effect mixing for 20 min. The resin was filtered, washed with DMF and resuspended in a solution of piperidine in DMF (30%, 10 mL). After gentle mixing for 40 min, the resin was filtered and washed with DMF, methylene chloride, methanol and methylene chloride and diluted with DMF (40 mL). To this mixture was added *N*-ethoxycarbonyl-D-thiazolidine-4-carboxylic acid (O-7, 0.15 g, 0.72 mmol), HOBt (0.11 g, 0.72 mmol), PyBOP (0.37 g, 0.72 mmol) and DIEA (313 μ L, 1.80 mmol). The reaction was mixed for 4 h at which point the qualitative Kaiser test was negative. The resin was filtered and washed with DMF, methylene chloride and MeOH and dried *in vacuo* to afford resin O-8. After swelling with a minimum of methylene chloride (ca. 0.5 mL), the resin O-8 was suspended with 95 % aqueous TFA (5 mL). The mixture was mixed by magnetic stirring for 1h, filtered and washed with TFA (2 X 3 mL) and methylene chloride. The combined filtrates were evaporated *in vacuo* to afford a residue that was purified by flash chromatography using methylene chloride/methanol (2%) containing glacial acetic acid (0.1%) as eluant to afford the title compound (80 mg). Lyophilization from glacial acetic acid afforded an amorphous powder: IR (drift) 3311, 3298, 1670, 1601, 1579, 1531, 1487, 1412, 1380, 1345, 1324, 1265, 1205, 1190, 709 cm^{-1} ; ^1H NMR (300 MHz, DMSO- d_6) δ 10.39 (1 H), 8.39 (1 H), 8.14 (2 H), 7.86 (2 H), 7.74 (3 H), 7.36 (2 H), 4.83 (2 H), 4.59 (1 H), 4.48 (1 H), 4.22 (2 H), 3.48 (2 H), 3.26 (1 H), 3.06 (2 H), 1.35 (3 H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 173.4, 169.8, 165.8, 154.0, 138.0, 135.4, 133.3, 131.9, 129.8, 128.8, 128.0, 120.4, 61.8, 54.2, 48.8, 36.7, 14.8; MS (ESI+) for $\text{C}_{23}\text{H}_{25}\text{N}_3\text{O}_6\text{S}$ m/z 472.0 ($\text{M}+\text{H}^+$); MS (ESI-) for $\text{C}_{23}\text{H}_{25}\text{N}_3\text{O}_6\text{S}$ m/z 470.1 ($\text{M}-\text{H}^-$); MS (FAB) m/z (rel. intensity) 472 (MH^+ , 99), 472 (99), 371 (31), 160 (31), 81 (31), 71 (45), 69 (46), 57 (71), 55 (58), 43 (47), 41 (42); HRMS (FAB) calcd for $\text{C}_{23}\text{H}_{25}\text{N}_3\text{O}_6\text{S} + \text{H}_1$ 472.1542, found 472.1563; Anal. Calcd for $\text{C}_{23}\text{H}_{25}\text{N}_3\text{O}_6\text{S} \cdot 1.5 \text{H}_2\text{O}$: C, 55.41; H, 5.66; N, 8.43. Found: C, 55.47; H, 5.21; N, 8.00.

Example 269.

(4S)-4-[[[(1S)-1-Carboxy-2-[4-[(acetyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester

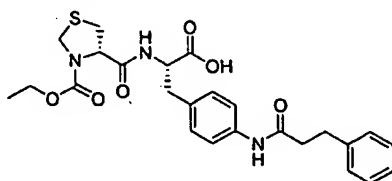
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The title compound was prepared as described in Scheme O using acetyl chloride to form the requisite amide. Physical data as follows: IR (drift) 3311, 1709, 1667, 1602, 1536, 1517, 1412, 1378, 1344, 1321, 1266, 1218, 1185, 1116, 769 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 9.79 (1 H), 8.15 (1 H), 7.37 (2 H), 7.02 (2 H), 4.54 (2 H), 4.31 (1 H), 4.19 (1 H), 3.95 (2 H), 3.12 (1 H), 2.93 (1 H), 2.75 (2 H), 1.82 (3 H), 1.09 (3 H); ^{13}C NMR (75 MHz, $\text{DMSO}-d_6$) δ 173.2, 169.8, 168.5, 154.0, 138.2, 132.4, 129.8, 119.1, 62.1, 61.7, 53.9, 36.7, 24.4, 21.5, 14.8; MS (ESI+) for $\text{C}_{18}\text{H}_{23}\text{N}_3\text{O}_6\text{S}$ m/z 410.0 ($\text{M}+\text{H}^+$); MS (ESI-) for $\text{C}_{18}\text{H}_{23}\text{N}_3\text{O}_6\text{S}$ m/z 408.0 ($\text{M}-\text{H}^-$); MS (FAB) m/z (rel. intensity) 410 (MH^+ , 99), 486 (20), 411 (22), 410 (99), 409 (9), 205 (22), 188 (9), 177 (9), 160 (35), 148 (9), 88 (14); HRMS (FAB) calcd for $\text{C}_{18}\text{H}_{23}\text{N}_3\text{O}_6\text{S} + \text{H}_1$ 410.1385, found 410.1379. Anal. Calcd for $\text{C}_{18}\text{H}_{23}\text{N}_3\text{O}_6\text{S} \cdot 0.3 \text{ H}_2\text{O}$: C, 52.11; H, 5.73; N, 10.13. Found: C, 51.73; H, 5.73; N, 9.82.

Example 270.

(4S)-4-[[[(1S)-1-Carboxy-2-[4-[(3-phenylpropanoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester



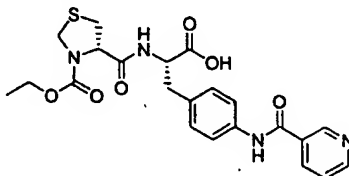
The title compound was prepared as described in Scheme O using hydrocinnamoyl chloride to form the requisite amide. Physical data as follows: IR (drift) 3311, 2978, 2930, 1665, 1601, 1534, 1517, 1413, 1379, 1344, 1252, 1216, 1187, 1115, 700 cm^{-1} ; ^1H NMR (300 MHz, $\text{CDCl}_3/\text{CD}_3\text{OD}$ (10%)) δ 7.34 (2 H), 7.21 (5 H), 7.02 (2 H), 4.67 (3 H), 4.30 (1 H), 4.09 (2 H), 3.11 (4 H), 2.97 (2 H), 2.58 (2 H), 1.19 (3 H); ^{13}C NMR (75 MHz, CDCl_3) δ 172.7, 171.3, 170.0, 154.8, 140.7, 137.0, 131.6, 129.7, 128.4, 128.2, 126.2, 119.9, 63.0, 62.6, 53.1, 38.9, 36.7, 31.5, 29.6, 14.2; MS (ESI+) for $\text{C}_{25}\text{H}_{29}\text{N}_3\text{O}_6\text{S}$ m/z 500.2

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(M+H)⁺; MS (ESI-) for C₂₅H₂₉N₃O₆S *m/z* 498.3 (M-H)⁻; Anal. Calcd for C₂₅H₂₉N₃O₆S: C, 60.10; H, 5.85; N, 8.41. Found: C, 59.85; H, 6.07; N, 8.09.

Example 271.

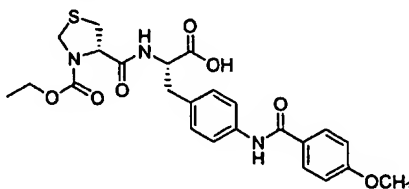
- 5 (4*S*)-4-[[[(1*S*)-1-Carboxy-2-[4-[(3-pyridinylcarbonyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester



- The title compound was prepared as described in Scheme O using nicotinoyl chloride to form the requisite amide. Physical data as follows: IR (drift) 3301, 3061, 2983, 2935, 1709, 1675, 1603, 1535, 1517, 1415, 1380, 1345, 1326, 1204, 1140 cm⁻¹; ¹H NMR (300 MHz, DMSO-*d*₆) δ 12.74 (1 H), 10.42 (1 H), 9.11 (1 H), 8.76 (1 H), 8.29 (2 H), 7.67 (2 H), 7.55 (1 H), 7.19 (2 H), 4.62 (2 H), 4.44 (1 H), 4.29 (1 H), 4.05 (2 H), 3.24 (1 H), 3.04 (1 H), 2.88 (2 H), 1.15 (3 H); ¹³C NMR (75 MHz, DMSO-*d*₆) δ 173.1, 170.0, 164.3, 154.0, 152.4, 149.0, 137.7, 135.9, 133.5, 131.1, 129.8, 123.9, 120.5, 62.4, 61.8, 53.8, 36.7, 14.8; MS (ESI+) for C₂₂H₂₄N₄O₆S *m/z* 473.3 (M+H)⁺; MS (ESI-) for C₂₂H₂₄N₄O₆S *m/z* 471.3 (M-H)⁻; HRMS (FAB) calcd for C₂₂H₂₄N₄O₆S +H₁ 473.1494, found 473.1509.

Example 272.

- 20 (4*S*)-4-[[[(1*S*)-1-Carboxy-2-[4-[(4-methoxybenzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester



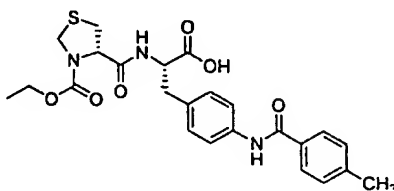
The title compound was prepared as described in Scheme O using *p*-anisoyl chloride to form the requisite amide. Physical data as follows: IR (drift) 1709, 1667,

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1604, 1532, 1514, 1439, 1412, 1379, 1343, 1323, 1255, 1221, 1178, 1027, 763 cm^{-1} ; ^1H NMR (300 MHz, $\text{CDCl}_3/\text{CD}_3\text{OD}$ (10%)) δ 7.79 (2 H), 7.47 (2 H), 7.05 (2 H), 6.87 (2 H), 4.68 (2 H), 4.58 (1 H), 4.28 (1 H), 4.07 (2 H), 3.79 (3 H), 3.66 (2 H), 3.04 (4 H), 1.17 (3 H); ^{13}C NMR (75 MHz, CDCl_3) δ 177.9, 174.0, 170.1, 166.3, 158.8, 141.1, 135.8, 133.7, 133.0, 130.8, 124.6, 117.7, 66.9, 66.6, 59.3, 40.7, 33.5, 24.5, 18.2; MS (ESI+) for $\text{C}_{24}\text{H}_{27}\text{N}_3\text{O}_7\text{S}$ m/z 502.0 ($\text{M}+\text{H}$) $^+$; MS (ESI-) for $\text{C}_{24}\text{H}_{27}\text{N}_3\text{O}_7\text{S}$ m/z 500.1 ($\text{M}-\text{H}$) $^-$; MS (FAB) m/z (rel. intensity) 502 (MH^+ , 52), 503 (17), 502 (52), 297 (12), 240 (12), 160 (21), 135 (99), 88 (12), 73 (20), 69 (13), 57 (12); HRMS (FAB) calcd for $\text{C}_{24}\text{H}_{27}\text{N}_3\text{O}_7\text{S} + \text{H}_1$ 502.1648, found 502.1657. Anal. Calcd for $\text{C}_{24}\text{H}_{27}\text{N}_3\text{O}_7\text{S} \cdot 0.3 \text{H}_2\text{O}$: C, 56.86; H, 5.49; N, 8.29. Found: C, 56.65; H, 5.34; N, 7.92.

Example 273.

(4S)-4-[[[(1S)-1-Carboxy-2-[4-[(4-methylbenzoyl)amino]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester

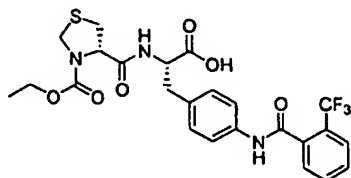


The title compound was prepared as described in Scheme O using *p*-tolouyl chloride to form the requisite amide. Physical data as follows: IR (drift) 3310, 2981, 2929, 1671, 1608, 1599, 1531, 1517, 1413, 1379, 1344, 1324, 1265, 1210, 1188 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 10.10 (1 H), 8.24 (1 H), 7.85 (2 H), 7.67 (2 H), 7.33 (2 H), 7.17 (2 H), 4.62 (2 H), 4.41 (1 H), 3.28 (2 H), 4.02 (2 H), 3.23 (1 H), 3.04 (1 H), 2.87 (2 H), 2.38 (3 H), 1.15 (3 H); ^{13}C NMR (75 MHz, $\text{DMSO}-d_6$) δ 173.2, 169.8, 165.6, 154.0, 141.9, 138.1, 133.1, 132.5, 129.7, 129.7, 128.1, 120.4, 62.4, 61.8, 54.0, 36.7, 21.4, 14.8; MS (ESI+) for $\text{C}_{24}\text{H}_{27}\text{N}_3\text{O}_6\text{S}$ m/z 486.2 ($\text{M}+\text{H}$) $^+$; HRMS (FAB) calcd for $\text{C}_{24}\text{H}_{27}\text{N}_3\text{O}_6\text{S} + \text{H}_1$ 486.1699, found 486.1713; Anal. Calcd for $\text{C}_{24}\text{H}_{27}\text{N}_3\text{O}_6\text{S} \cdot 0.3 \text{H}_2\text{O}$: C, 58.71; H, 5.67; N, 8.56. Found: C, 58.37; H, 5.67; N, 8.35.

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Example 274.

(4S)-4-[[[(1S)-1-Carboxy-2-[4-[[2-(trifluoromethyl)benzoyl]amino]phenyl]ethyl]amino]-carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester



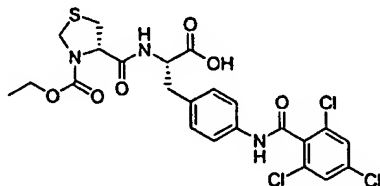
5

The title compound was prepared as described in Scheme O using (2-trifluoromethyl)benzoyl chloride to form the requisite amide. Physical data as follows:
 IR (drift) 3295, 1709, 1663, 1603, 1533, 1518, 1414, 1380, 1344, 1316, 1269, 1176, 1132,
 1108, 769 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 10.48 (1 H), 8.23 (1 H), 7.77 (2 H),
 7.58 (2 H), 7.17 (2 H), 4.64 (2 H), 4.44 (1 H), 4.29 (1 H), 3.99 (2 H), 3.24 (1 H), 3.05 (2
 H), 1.09 (3 H); ^{13}C NMR (75 MHz, $\text{DMSO}-d_6$) δ 173.2, 169.2, 165.8, 154.1, 137.8, 136.7,
 133.0, 130.4, 129.9, 128.9, 126.7, 126.4, 126.0, 119.8, 62.2, 61.8, 54.0, 37.1, 14.7; MS
 (ESI+) for $\text{C}_{24}\text{H}_{25}\text{F}_3\text{N}_3\text{O}_6\text{S}$ m/z 540.0 ($\text{M}+\text{H}^+$); MS (ESI-) for $\text{C}_{24}\text{H}_{25}\text{F}_3\text{N}_3\text{O}_6\text{S}$ m/z 538.1 ($\text{M}-$
 H); HRMS (FAB) calcd for $\text{C}_{24}\text{H}_{24}\text{F}_3\text{N}_3\text{O}_6\text{S} + \text{H}$, 540.1416, found 540.1423; Anal. Calcd
 for $\text{C}_{24}\text{H}_{24}\text{F}_3\text{N}_3\text{O}_6\text{S} \cdot 0.5 \text{H}_2\text{O}$: C, 51.49; H, 4.70; N, 7.83. Found: C, 51.42; H, 4.42; N,
 7.45.

20

Example 275.

(4S)-4-[[[(1S)-1-Carboxy-2-[4-[(2,4,6-trichlorobenzoyl)amino]phenyl]ethyl]amino]-carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester

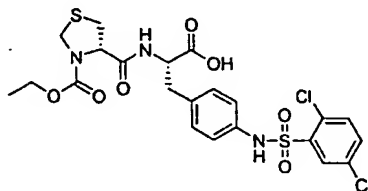


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The title compound was prepared as described in Scheme O using 2,4,6-trichlorobenzoyl chloride to form the requisite amide. Physical data as follows: IR (drift) 3286, 2926, 1709, 1664, 1604, 1578, 1542, 1517, 1413, 1379, 1345, 1325, 1269, 1218, 1187 cm^{-1} ; ^1H NMR (300 MHz, $\text{CDCl}_3/\text{CD}_3\text{OD}$ (10%)) δ 7.58 (2 H), 7.41 (2 H), 7.20 (2 H), 4.84 (2 H), 4.69 (1 H), 4.41 (1 H), 4.19 (2 H), 3.27 (4 H), 1.26 (3 H); ^{13}C NMR (75 MHz, $\text{DMSO}-d_6$) δ 173.0, 169.8, 161.5, 154.0, 137.2, 135.9, 135.1, 133.9, 132.6, 130.1, 128.5, 119.6, 62.2, 61.7, 54.0, 36.7, 14.8; MS (FAB) m/z (rel. intensity) 574 (MH^+ , 95), 576 (96), 574 (95), 160 (99), 91 (79), 88 (40), 69 (64), 57 (59), 55 (59), 43 (56), 41 (39); HRMS (FAB) calcd for $\text{C}_{23}\text{H}_{22}\text{Cl}_3\text{N}_3\text{O}_6\text{S} + \text{H}_1$ 574.0373, found 574.0364. Anal. Calcd for $\text{C}_{23}\text{H}_{22}\text{Cl}_3\text{N}_3\text{O}_6\text{S}$: C, 48.06; H, 3.86; N, 7.31; Cl, 18.50. Found: C, 48.52; H, 4.13; N, 7.08.

Example 276.

(4S)-4-[[[(1S)-1-Carboxy-2-[4-[(2,5-dichlorophenyl)sulfonyl]amino] phenyl]ethyl]-amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester



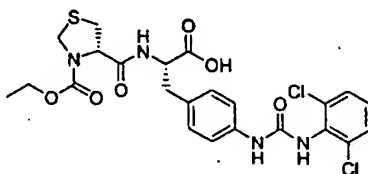
The title compound was prepared as described in Scheme O using 2,5-dichlorobenzene sulfonyl chloride to form the requisite sulfonamide. Physical data as follows: IR (drift) 1709, 1676, 1531, 1512, 1450, 1428, 1409, 1378, 1344, 1221, 1167, 1143, 1113, 1101, 1041 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 8.36 (1 H), 7.90 (3 H), 7.20 (2 H), 7.12 (2 H), 4.70 (2 H), 4.47 (1 H), 4.34 (1 H), 4.10 (2 H), 3.19 (4 H), 1.27 (3 H); ^{13}C NMR (75 MHz, $\text{DMSO}-d_6$) δ 173.0, 169.8, 153.9, 138.7, 135.5, 134.8, 134.1, 132.6, 131.0, 130.5, 130.0, 120.0, 62.4, 61.7, 53.5, 36.3, 14.8; MS (FAB) m/z (rel. intensity) 576 (MH^+ , 99), 652 (27), 578 (83), 577 (31), 576 (99), 160 (98), 106 (47), 88 (40), 81 (32), 69 (31), 57 (28); HRMS (FAB) calcd for $\text{C}_{22}\text{H}_{23}\text{Cl}_2\text{N}_3\text{O}_7\text{S}_2 + \text{H}_1$ 576.0433,

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found 576.0400. Anal. Calcd for $C_{22}H_{23}Cl_2N_3O_7S_2 \cdot 0.1 H_2O$: C, 45.70; H, 4.04; N, 7.27.
Found: C, 45.94; H, 4.04; N, 6.87.

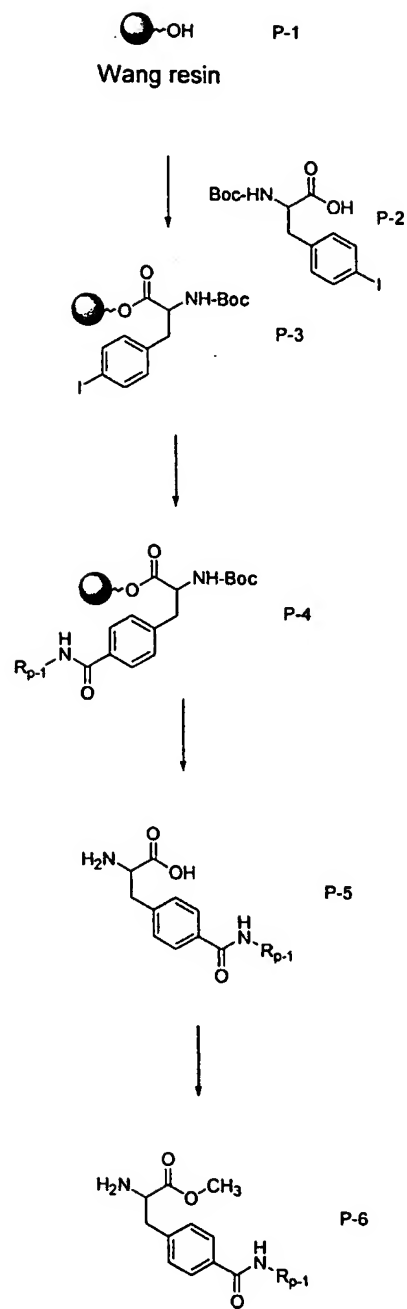
Example 277.

- 5 (4S)-4-[[[(1S)-1-Carboxy-2-[4-[[[(2,6-dichlorophenyl)amino]carbonyl] amino]phenyl]-ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester

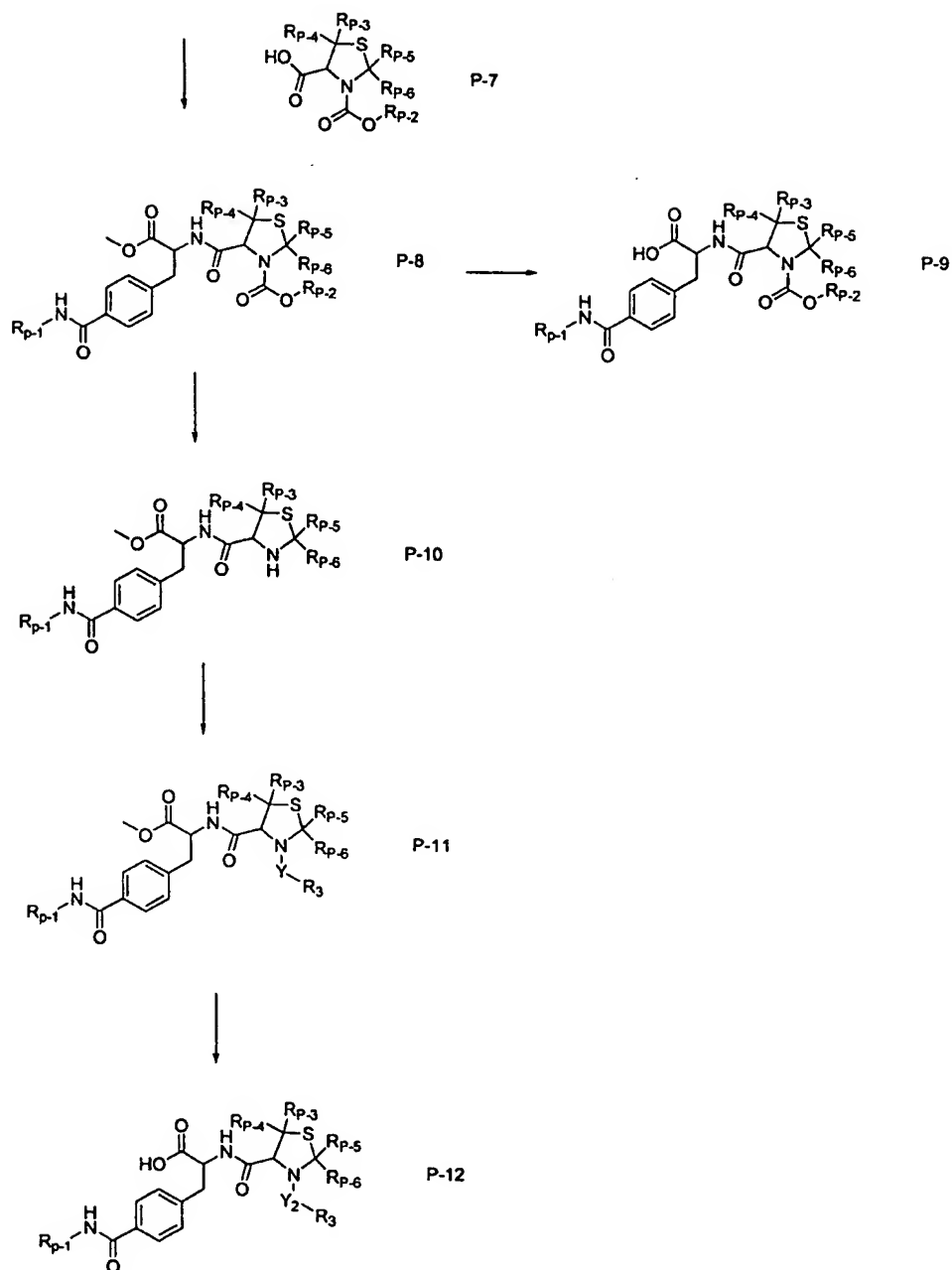


- 10 The title compound was prepared as described in Scheme O using 2,6-dichlorophenyl isocyanate to form the requisite urea. Physical data as follows: IR (drift) 3284, 3277, 1709, 1655, 1600, 1569, 1544, 1452, 1431, 1415, 1347, 1238, 1217, 1195, 771 cm^{-1} ; 1H NMR (300 MHz, DMSO- d_6) δ 9.09 (1 H), 8.41 (1 H), 8.09 (1 H), 7.52 (2 H), 7.31 (3 H), 7.08 (2 H), 4.62 (2 H), 4.35 (1 H), 4.28 (1 H), 3.27 (1 H), 3.02 (1 H), 2.85 (2 H), 1.10 (3 H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 173.6, 169.8, 154.0, 152.8, 138.7, 134.4, 133.9, 131.3, 130.0, 128.8, 128.7, 118.1, 62.4, 61.8, 54.3, 36.7, 14.8; HRMS (FAB) calcd for $C_{23}H_{24}Cl_2N_4O_6S + H_1$, 555.0872, found 555.0877. Anal. Calcd for $C_{23}H_{24}Cl_2N_4O_6S \cdot 2 H_2O$: C, 46.71; H, 4.77; N, 9.47. Found: C, 47.08; H, 4.53; N, 9.06.
- 15

Scheme P



Scheme P (continued)

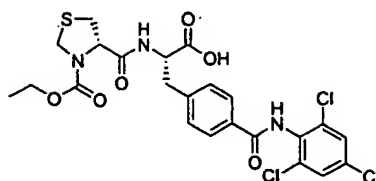


Where: $R_{p,1}$ is defined as $R_{1,2}$; $R_{p,2}$ is defined as C_{1-6} alkyl or C_{7-17} arylalkyl; $R_{p,3}$, $R_{p,4}$, and $R_{p,5}$, are defined independently as R_1 . $R_{p,6}$ is defined as R_2 .

Scheme P describes a method for the preparation of examples of the formula **P-9** and **P-12**. Commercially available Wang resin (**P-1**) is acylated with commercially available *N*- α -Boc-Phe(I)-OH (**P-2**) under standard conditions to afford the resin of formula **P-3**. Carbonylation of the resin bound aryl iodide with carbon monoxide and an amine in the presence of a source of palladium(0) affords the resin bound amide of general formula **P-5** (for a general review of carbonylation chemistry, see: Colquhoun, H.M.; Thompson, D.J.; Twigg, M.V. *Carbonylation* Plenum Press: New York, 1991). Mild cleavage under standard conditions affords the amino acid of general structure **P-5** which is esterified under mild acid catalysis to afford the amino ester of general structure **P-6**. Condensation with a commercially available or readily prepared thiazolidine-4-carboxylic acid of general formula **P-7** under conditions described in Scheme A affords the pseudodipeptide of general structure **P-8**. Mild base hydrolysis of the ester of general structure **P-8** affords the acid of general structure **P-9**. In the case where $R_{p,2}$ is a 9-fluorenylmethyl group, standard Fmoc group removal (Atherton, E.; Sheppard R.C. *Solid Phase Peptide Synthesis: A Practical Approach*; IRL Press at Oxford University Press: Oxford, 1989) affords the amine of general structure **P-10**, which may be reacted with a variety of electrophilic reagents as described in Scheme A to afford amides, ureas, sulfonamides and carbamates of general structure **P-11**. Mild base hydrolysis of the ester in general structure **P-11** affords the acid of general structure **P-12**.

Preparation 80 and Example 278.

(4*S*)-4-[[[(1*S*)-1-Carboxy-2-[4-[(2,4,6-trichlorophenyl)amino]carbonyl] phenyl]ethyl]-amino]carbonyl]-3-thiazolidinecarboxylic acid 3-ethyl ester



To a cooled (0-50°C) mixture of Wang polystyrene resin P-1 (Advanced Chemtech, 2.0 g, ca. 1.5 mmol), *N*-Boc-4-iodo-L-phenylalanine P-2 (Bachem, 4.00 g, 10 mmol), and PPh₃ (1.30 g, 5.0 mmol) in THF (20 mL) was added diethyl azodicarboxylate (0.80 mL, 5.0 mmol) in 4 approximately equal portions at 5 min intervals. When the orange color had discharged, the mixture was warmed to ambient temperature and stirred for 5 h. The mixture was diluted with THF (30 mL) and filtered. The resin was washed with DMF, THF and MeOH and dried *in vacuo* to afford the esterified resin P-3 (2.68 g) as a colorless powder: ¹³C NMR (100 MHz, CD₂Cl₂, 4 mm MAS probe) δ 171.86, 155.33, 137.85, 136.40, 131.87, 128.00, 92.74, 80.09, 54.05, 38.05, 28.51.

N₂ was bubbled through a mixture of *N*-Boc-4-iodo-L-phenylalanine functionalized Wang resin P-3 (500 mg, ca. 0.3 mmol), PPh₃ (0.21 g, 0.8 mmol), 2,4,6-trichloroaniline (0.98 g, 5.0 mmol) and DIEA (3.48 mL, 20 mmol) in NMP (20 mL) for 10 min. Pd₂dba₃ (0.18 g, 0.2 mmol) was added and the reaction mixture was placed under a CO atmosphere and heated (bath temp. 70 °C) for 18 h. Upon cooling to ambient temperature, the mixture was diluted with 3% (w/v) sodium diethyldithiocarbamate in 95:5 NMP:DIEA (10 mL). After an additional 10 min, the mixture was filtered and the resin washed with NMP, THF and MeOH and dried *in vacuo* to afford the functionalized resin P-4 as a colorless powder.

Resin P-4 was swollen with methylene chloride (0.5 mL) and diluted with 95:5 TFA:H₂O (10 mL). After 90 min the mixture was filtered and the resin washed with TFA (3 x 5 mL) and CH₂Cl₂. The combined filtrates were concentrated *in vacuo* and the residue lyophilized from glacial acetic acid to provide the amino acid P-5 (152 mg) as a powder which was used without purification. Physical data as follows: MS (FAB) *m/z* (rel. intensity) 387 (M+H, 42), 427 (26), 426 (80), 389 (46), 387 (42), 366 (33), 279 (99), 177 (54), 146 (18), 119 (26), 23 (26); HRMS (FAB) calcd for C₁₆H₁₃Cl₃N₂O₃+H₁, 387.0070, found 387.0084.

The amino acid P-5 was dissolved in methanolic HCl (20 mL) and stirred at room temperature for 18 h. Concentration *in vacuo* afforded the methyl ester P-6 which was used without purification. Physical data as follows: MS (ES+) for $C_{17}H_{15}Cl_3N_2O_3$ m/z 400.9 (M+H)⁺.

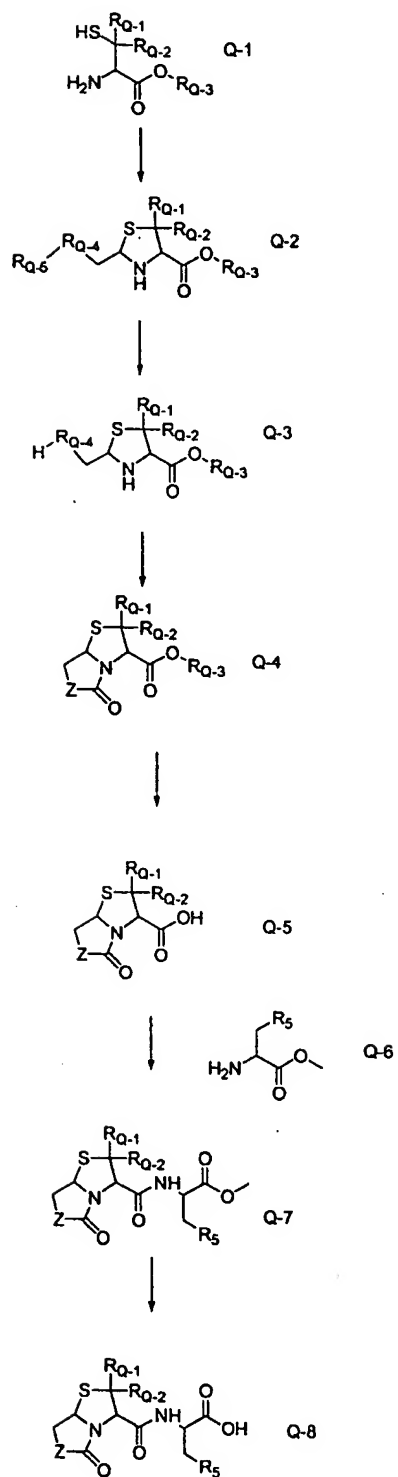
- 5 To a cooled (0-5 °C) solution of *N*-ethoxycarbonyl-D-thiazolidine-4-carboxylic acid P-7 (82 mg, 0.4 mmol) and HOAt (54 mg, 0.4 mmol) in CH_2Cl_2 /DMF (1:1, 4 mL) was added EDC (76 mg, 0.4 mmol). After stirring for 10 min, the solution was added to the amino ester P-6 described above at 0-5 °C followed by DIEA (208 μ L, 1.2 mmol). After an additional 30 min at 0-5 °C, the solution was allowed to warm to room
- 10 temperature and stirred an additional hour. Volatiles were removed *in vacuo* and the residue partitioned between ethyl acetate and 0.1 N aqueous HCl. The organic layer was separated, washed with 0.1 N aqueous HCl, saturated aqueous $NaHCO_3$, brine, dried ($MgSO_4$), filtered and concentrated *in vacuo*. Purification of the residue by flash chromatography using CH_2Cl_2 /ethyl acetate/hexanes (1:1:2) containing 2-propanol (0.1 %) as eluant afforded the ester P-8 as a powder: ¹H NMR (300 MHz, $CDCl_3$) δ 7.99 (1 H), 7.87 (2 H), 7.40 (2 H), 7.24 (2 H), 4.92 (1 H), 4.70 (2 H), 4.34 (1 H), 4.10 (2 H), 3.74 (3 H), 3.20 (4 H), 1.25 (3 H); ¹³C NMR (75 MHz, $CDCl_3$) δ 171.2, 169.7, 165.4, 154.9, 140.6, 134.3, 133.4, 132.2, 131.3, 129.7, 128.4, 127.9, 63.1, 62.7, 60.4, 53.0, 52.6, 37.6, 21.0, 14.5; MS (ESI+) for $C_{24}H_{24}Cl_3N_3O_6S$ m/z 589.9 (M+H)⁺; MS (ESI-) for
- 15 $C_{24}H_{24}Cl_3N_3O_6S$ m/z 588.0 (M-H)⁻.
- 20

- To a cooled (0-5°C) solution of the ester P-8 (72 mg, 0.12 mmol) in THF (5 mL) and water (0.5 mL) was added an 0.1 N aqueous solution of NaOH (1.3 mL, 0.13 mmol) via a syringe pump over 1 h. After an additional 45 min at 0-5°C, the reaction mixture was diluted with ethyl acetate and acidified with 0.25 N HCl to a pH of ca. 3. The organic
- 25 layer was separated, washed with water and concentrated *in vacuo*. Purification of the residue by flash chromatography using methylene chloride and methanol (0-5%) as eluant provided a solid which was crystallized from ethyl acetate/ CH_2Cl_2 /hexanes to afford the title compound (45 mg) as colorless solid: IR (drift) 1743, 1726, 1709, 1691, 1675, 1663, 1553, 1521, 1490, 1428, 1415, 1379, 1345, 1290, 1189 cm^{-1} ; ¹H NMR (300 MHz, DMSO-*d*₆) δ 10.27 (1 H), 8.35 (1 H), 7.92 (, 2 H), 7.81 (2 H), 7.37 (2 H), 4.62 (2 H), 4.53 (1 H),
- 30

4.29 (2 H), 4.00 (2 H), 3.11 (3 H), 2.77 (1 H), 1.12 (3 H); ^{13}C NMR (75 MHz, DMSO- d_6) δ 172.9, 170.0, 165.4, 154.0, 142.5, 135.5, 133.9, 133.0, 131.6, 129.8, 128.7, 128.0, 62.0, 61.7, 53.4, 37.0, 14.8; MS (FAB) m/z (rel. intensity) 574 (MH^+ , 80), 576 (80), 574 (80), 379 (99), 160 (82), 91 (95), 81 (72), 69 (93), 57 (86), 55 (86), 43 (90); HRMS (FAB) calcd
5 for $\text{C}_{23}\text{H}_{22}\text{Cl}_3\text{N}_3\text{O}_6\text{S} + \text{H}$, 574.0373, found 574.0358.

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Scheme Q



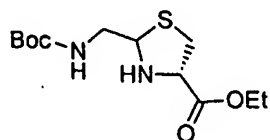
R_{Q-1} and R_{Q-2} are defined independently as R_1 . R_{Q-3} is defined as C_{1-6} alkyl or C_{7-17} arylalkyl. R_{Q-4} is defined as oxygen or $N-R_{11}$. R_{Q-5} is defined as a suitable protecting group for a nitrogen (such as Boc or Fmoc) or oxygen (such as t-butyldimethylsilyl) (Greene, T.W.; Wuts, P.G.M.. *Protective Groups in Organic Synthesis*, John Wiley and Sons, New York, 1991).

Scheme Q describes a general method for the preparation of examples of the formula Q-7. A commercially available or readily prepared sulfur containing amino acid of structure Q-1 is condensed with a suitably protected aldehyde to afford the thiazolidine-4-carboxylic acid of general formula Q-2. Standard deprotection (Greene, T.W.; Wuts, P.G.M.. *Protective Groups in Organic Synthesis*, John Wiley and Sons, New York, 1991) affords intermediate Q-3 which is readily cyclized to the bicycle Q-4 using 1,1'-carbonyldiimidazole or phosgene or a suitable equivalent. For the preparation of bicycles of general structure Q-4 in which Z is CH_2 , see as examples: (a) Aszodi, J.; Bonnet, A.; Teutsch, G. *Tetrahedron* **1990**, *46*, 1579. (b) Baldwin, J.E.; Lee, V.; Schofield, C.J. *Heterocycles* **1992**, *34*, 903. (c) Genin, M.J.; Johnson, R.L. *J. Am. Chem. Soc.* **1992**, *114*, 8778. (d) Siddiqui, M.A.; Preville, P.; Tarazi, M.; Warder, S.E.; Eby, P.; Gorseth, E.; Puumala, K.; DiMaio, J. *Tetrahedron Lett.* **1997**, *38*, 8807. (e) Subasinghe, M.L.; Bontems, R.J.; McIntee, E.; Mishra, R.K.; Johnson, R.L. *J. Med. Chem.* **1993**, *36*, 2356. Removal of the ester protecting group affords the acid of general structure Q-5 which is condensed with amino acyl derivative Q-6 under standard peptide synthesis conditions to provide Q-7 (for a review of procedures of peptide synthesis see: Bodansky, M.; Bodansky, A. *The Practice of Peptide Synthesis*; Springer-Verlag: Berlin, 1984). Mild base hydrolysis of the ester of general structure Q-7 provides acid Q-8.

25

Preparation 81.

(Scheme Q, Q-2: where R_{Q-1} and R_{Q-2} are equal to hydrogen, R_{Q-3} is ethyl, R_{Q-4} is NH, R_{Q-5} is Boc and stereochemistry is (S)).



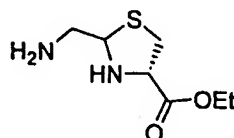
-233-

To a suspension of D-cysteine 1.5 hydrate hydrochloride (Q-1, Scheme Q where R_{Q-1} and R_{Q-1} are equal to hydrogen, R_{Q-3} is hydrogen and stereochemistry is (S)) (5 g, 27.1 mmol) in absolute ethyl alcohol (50 mL) was added triethylorthoformate (13.5 mL, 81.2 mmol) at ambient temperature. A stream of anhydrous HCl gas was bubbled through the solution for 30 min. The stream of anhydrous HCl gas was maintained as the mixture heated to 70°C for 2 h. The reaction mixture was concentrated *in vacuo* and the resulting residue triturated with diethyl ether to afford D-cysteine ethyl ester (4.43 g) as a white solid which was used without further purification. Physical data as follows: ^1H NMR (300 MHz, D_2O) δ 4.36 (1 H), 4.27 (2 H), 3.12 (2 H), 1.25 (3 H); MS (ESI+) for $\text{C}_5\text{H}_{11}\text{NO}_2\text{S}$ m/z 150.0 (M+H) $^+$.

To a solution of D-cysteine ethyl ester (1.89 g, 10.2 mmol) in H_2O (46 mL) was added potassium acetate (1.22 g, 12.4 mmol) and *t*-butyl *N*-(2-oxoethyl)carbamate (Aldrich, 2.38 g, 12.0 mmol based on 80% purity as determined by ^1H NMR) in ethyl alcohol (46 mL) at ambient temperature. After 8 h, the reaction mixture was concentrated *in vacuo* and the residue purified by flash chromatography using methylene chloride/methanol (1%) as eluant to afford the title compound (1.9 g) as an oil: ^1H NMR (300 MHz, CDCl_3) δ 5.01 (1 H), 4.75 (1 H), 4.26 (2 H), 3.93 (1 H), 3.37 (2 H), 3.12 (1 H), 2.91 (1 H), 1.47 (9 H), 1.32 (3 H); MS (ESI+) for $\text{C}_{12}\text{H}_{22}\text{N}_2\text{O}_4\text{S}$ m/z 291.1 (M+H) $^+$. MS (ESI-) for $\text{C}_{12}\text{H}_{22}\text{N}_2\text{O}_4\text{S}$ m/z 288.9 (M-H) $^-$.

Preparation 82.

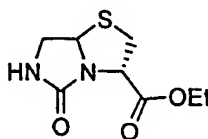
(Scheme Q, Q-3: where R_{Q-1} and R_{Q-1} are equal to hydrogen, R_{Q-3} is ethyl, R_{Q-4} is NH, and stereochemistry is (S)).



To a cooled (10-15°C) solution of Q-2 (Scheme Q where R_{Q-1} and R_{Q-1} are equal to hydrogen, R_{Q-3} is ethyl, R_{Q-4} is NH, R_{Q-5} is Boc and stereochemistry is (S)) (1.9 g) in dioxane (38 mL) was added dropwise anhydrous 4 M HCl in dioxane (156 mL). The solution was allowed to warm to ambient temperature and stirred for 2 h. The reaction mixture was concentrated *in vacuo* and azeotrope three times with methanol which afforded the title compound (1.72 g) as a tacky yellow solid: ¹H NMR (300 MHz, CDCl₃) δ 8.57 (2 H), 5.80 (1 H), 5.25 (1 H), 4.39 (2 H), 4.17 (1 H), 3.79 (4 H), 1.37 (3 H); MS (ESI+) for C₇H₁₄N₂O₂S *m/z* 191.1 (M+H)⁺. MS (ESI-) for C₇H₁₄N₂O₂S • 2HCl *m/z* 261.0 (M-H).

Preparation 83.

(Scheme Q, Q-4: where R_{Q-1} and R_{Q-1} are equal to hydrogen, R_{Q-3} is ethyl, Z is NH, and stereochemistry is (S)).



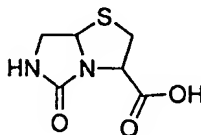
To a cooled (0-5°C) solution of Q-3 (Scheme Q where R_{Q-1} and R_{Q-1} are equal to hydrogen, R_{Q-3} is ethyl, R_{Q-4} is NH, and stereochemistry is (S)) (1.72 g, 6.54 mmol) in THF (650 mL) was added triethylamine (2.83 mL, 20.3 mmol) and 1,1'-carbonyldiimidazole (1.11 g, 6.87 mmol). After 3 d at ambient temperature, the mixture was recooled (0-5°C), treated with additional 1,1'-carbonyldiimidazole (530 mg, 3.27 mmol) and allowed to warm to ambient temperature. After 18 h, the reaction mixture was concentrated *in vacuo* and the resulting residue was partitioned between ethyl acetate and 0.25 N HCl. The organic layer was separated, washed with brine, dried (MgSO₄), filtered and concentrated *in vacuo*. The diastereomeric mixture was separated by chiral chromatography [5 x 25 cm (R,R) Whelk-O I, 50 mL/min 40% Isopropanol/heptane, 210 nm,]. Further purification of each isolated diastereomer by flash chromatography using methylene chloride/ethyl acetate (25%) as eluant afforded the diastereomers (255 mg, 464 mg) as oils. Physical data for the faster eluting diastereomer (analytical column conditions 0.46 x 25 cm (R,R)

Whekk-O I, 0.5 mL/min 40% IPA/heptane, 210 nm) as follows: ^1H NMR (300 MHz, CDCl_3) δ 5.51 (1 H), 5.15 (1 H), 4.32 (2 H), 3.96 (1 H), 3.86 (1 H), 3.53 (1 H), 3.26 (1 H), 3.13 (1 H), 1.34 (3 H); MS (ESI+) for $\text{C}_8\text{H}_{12}\text{N}_2\text{O}_3\text{S}$ m/z 217.1 ($\text{M}+\text{H}$) $^+$. MS (ESI+) for $\text{C}_8\text{H}_{12}\text{N}_2\text{O}_3\text{S}$ m/z 239.0 ($\text{M}+\text{Na}$) $^+$. MS (ESI-) for $\text{C}_8\text{H}_{12}\text{N}_2\text{O}_3\text{S}$ m/z 215.1 ($\text{M}-\text{H}$) $^-$.

5

Preparation 84.

(Scheme Q, Q-5: where $\text{R}_{\text{Q-1}}$ and $\text{R}_{\text{Q-1}}$ are equal to hydrogen, and Z is NH).



10 To a cooled (0-5°C) solution of the faster eluting diastereomer of general structure Q-4 (Scheme Q where $\text{R}_{\text{Q-1}}$ and $\text{R}_{\text{Q-1}}$ are equal to hydrogen, $\text{R}_{\text{Q-3}}$ is ethyl, Z is NH, and stereochemistry is (S)) (100 mg, 0.46 mmol) in THF (13 mL) and H_2O (1.5 mL) was added via syringe pump over 1 h 0.1 N NaOH (9.7 mL, 0.97 mmol). The reaction mixture was stirred for 2 h at 0°C, acidified with 1.0 N HCl (0.97 mL) and concentrated *in vacuo*. The

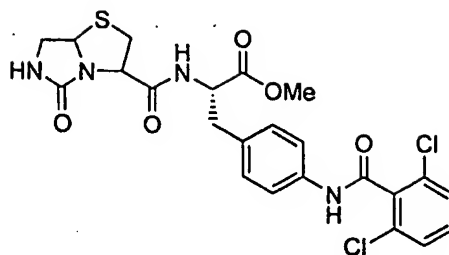
15 resulting residue was dried over P_2O_5 in a vacuum desiccator to afford the title compound (87 mg) as a glassy solid which was used without further purification: MS (ESI+) for $\text{C}_6\text{H}_8\text{N}_2\text{O}_3\text{S}$ m/z 189.0 ($\text{M}+\text{H}$) $^+$. MS (ESI+) for $\text{C}_6\text{H}_8\text{N}_2\text{O}_3\text{S}$ m/z 211.0 ($\text{M}+\text{Na}$) $^+$. MS (ESI-) for $\text{C}_6\text{H}_8\text{N}_2\text{O}_3\text{S}$ m/z 187.0 ($\text{M}-\text{H}$) $^-$.

20

Preparation 85.

(Scheme Q, Q-7: where $\text{R}_{\text{Q-1}}$ and $\text{R}_{\text{Q-1}}$ are equal to hydrogen, Z is NH, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry of the C-terminal amino acid is (S)).

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To a cooled (0-5°C) suspension of **Q-5** (Scheme Q where R_{Q-1} and R_{Q-1} are equal to hydrogen, and Z is NH) (87 mg, 0.46 mmol) in methylene chloride (10 mL) was added *O*-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate (HATU) (175 mg, 0.46 mmol), **Q-6** (Scheme Q where R_s is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry is (*S*)) (204 mg, 0.51 mmol) and *N,N*-diisopropylethylamine (0.24 mL, 1.38 mmol). After 7 h, the reaction mixture was diluted with methylene chloride, washed with 0.1 N HCl and brine, dried ($MgSO_4$), filtered and concentrated *in vacuo*. Purification of the residue by flash chromatography using methylene chloride/ethyl acetate (25%) as eluant afforded the title compound (89 mg) as a white solid: 1H NMR (300 MHz, $CDCl_3$) δ 7.57 (2 H), 7.31 (3 H), 7.13 (2 H), 4.97 (1 H), 4.82 (1 H), 4.47 (1 H), 3.80 (4 H), 3.55 (1 H), 3.46 (1 H), 3.34 (1 H), 3.08 (2 H); MS (ESI+) for $C_{23}H_{22}Cl_2N_4O_5S$ m/z 537.0 ($M+H$)⁺. MS (ESI+) for $C_{23}H_{22}Cl_2N_4O_5S$ m/z 558.9 ($M+Na$)⁺.

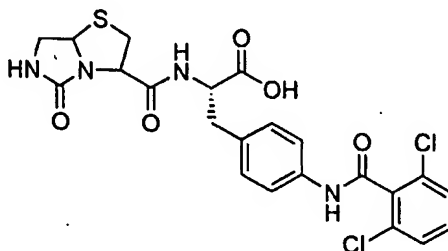
15

Preparation 86 and Example 279.

4-[(2,6-Dichlorobenzoyl)amino]-*N*-[[[(7*S*)-hexahydro-5-oxoimidazo[5,1-*b*]thiazol-3-yl]-carbonyl]-*L*-phenylalanine

(Scheme Q, **Q-8**: where R_{Q-1} and R_{Q-1} are equal to hydrogen, Z is NH, R_s is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry of the C-terminal amino acid is (*S*)).

20



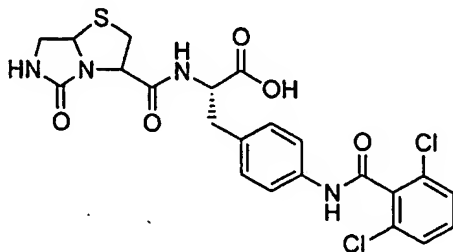
-237-

To a cooled (0-5°C) solution of Q-7 (Scheme Q where R_{Q-1} and R_{Q-1} are equal to hydrogen, Z is NH, R_s is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry of the C-terminal amino acid is (S)) (88 mg, 0.16 mmol) in THF (5 mL) and H₂O (0.5 mL) was added via syringe pump over 1 h 0.1 N NaOH (3.4 mL, 0.34 mmol). After 2 h, the reaction mixture was partitioned between ethyl acetate and 0.1 N HCl (7 mL) and diluted with H₂O (20 mL). The organic layer was separated, washed with H₂O, dried (MgSO₄), filtered and concentrated *in vacuo*. The resulting white solid was lyophilized from glacial acetic acid to afford the title compound (22 mg) as white solid: IR (drift) 2924, 1726, 1720, 1663, 1657, 1608, 1515, 1456, 1431, 1402, 1398, 1243, 1194, 797, 780 cm⁻¹; ¹H NMR (300 MHz, CD₃OD) δ 8.05 (1 H), 7.61 (2 H), 7.46 (3 H), 7.25 (2 H), 4.93 (2 H), 4.74 (2 H), 4.47 (1 H), 3.77 (1 H), 3.39 (2 H), 3.25 (1 H), 3.04 (2 H); ¹³C NMR (75 MHz, CD₃OD) δ 171.8, 165.2, 164.7, 138.2, 137.7, 135.1, 133.4, 132.4, 130.9, 129.4, 121.7, 66.3, 64.4, 54.5, 44.9, 37.4, 33.9; HRMS (FAB) calcd for C₂₂H₂₀Cl₂N₄O₅S + H, 523.0610, found 523.0629. MS (ESI+) for C₂₂H₂₀Cl₂N₄O₅S *m/z* 523.0 (M+H)⁺. MS (ESI-) for C₂₂H₂₀Cl₂N₄O₅S *m/z* 521.1 (M-H)⁻.

Example 280.

4-[(2,6-Dichlorobenzoyl)amino]-N-[[[(7aR)-hexahydro-5-oxoimidazo[5,1-b]thiazol-3-yl]-carbonyl]-L-phenylalanine

(Scheme Q, Q-8: where R_{Q-1} and R_{Q-1} are equal to hydrogen, Z is NH, R_s is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry of the C-terminal amino acid is (S)).



Example 280 was prepared as described in Scheme Q from the slower eluting diastereomer of general structure Q-4 (Scheme Q where R_{Q-1} and R_{Q-1} are equal to hydrogen, R_{Q-3} is ethyl, and Z is NH). Physical data as follows: IR (drift) 3251, 3079, 1730, 1662, 1611, 1549, 1516, 1482, 1431, 1333, 1306, 1269, 1229, 1196, 792 cm^{-1} ; ^1H NMR (300 MHz, CD_3OD) δ 7.59 (2 H), 7.44 (3 H), 7.24 (2 H), 4.65 (1 H), 3.81 (1 H), 3.44 (1 H), 3.23 (2 H), 3.08 (2 H); ^{13}C NMR (75 MHz, CD_3OD) δ 175.3, 172.22, 164.6, 155.3, 139.7, 139.1, 136.1, 134.1, 133.9, 132.3, 131.8, 130.9, 122.0, 68.4, 64.6, 45.6, 44.0, 36.3; HRMS (FAB) calcd for $\text{C}_{22}\text{H}_{20}\text{Cl}_2\text{N}_4\text{O}_5\text{S} + \text{H}_1$ 523.0610, found 523.0629. MS (ESI+) for $\text{C}_{22}\text{H}_{20}\text{Cl}_2\text{N}_4\text{O}_5\text{S}$ m/z 523.0 (M+H) $^+$. MS (ESI-) for $\text{C}_{22}\text{H}_{20}\text{Cl}_2\text{N}_4\text{O}_5\text{S}$ m/z 521.0 (M-H) $^-$. Anal. Calcd for $\text{C}_{22}\text{H}_{20}\text{Cl}_2\text{N}_4\text{O}_5\text{S} \cdot 0.13 \text{ H}_2\text{O}$: C, 50.26; H, 3.88; N, 10.66. Found: C, 50.72; H, 3.96; N, 10.13. % Water (KF): 0.45.

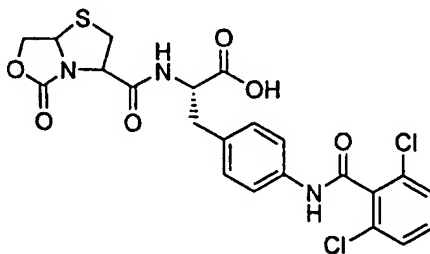
15

Example 281.

4-[(2,6-Dichlorobenzoyl)amino]-*N*-[(tetrahydro-5-oxo-5*H*-thiazolo[3,2-*c*]oxazol-3-yl)-carbonyl]-L-phenylalanine (less polar diastereomer)

(Scheme Q, Q-8: where R_{Q-1} and R_{Q-1} are equal to hydrogen, Z is O, R_s is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry of the C-terminal amino acid is (*S*)).

20



Example 281 was prepared as described in Scheme Q using commercially available (*t*-butyldimethylsilyloxy)acetaldehyde to form the requisite thiazolidine-4-carboxylic acid.

Physical data as follows: IR (drift) 3293, 3194, 1754, 1667, 1603, 1533, 1517, 1431,

1411, 1392, 1324, 1266, 1206, 798, 781 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 8.44 (1

5 H), 7.53 (5 H), 7.21 (2 H), 5.00 (1 H), 4.84 (1 H), 4.65 (1 H), 4.42 (2 H), 3.08 (2 H), 2.91

(1 H); ^{13}C NMR (75 MHz, CD_3OD) δ 169.4, 163.8, 160.6, 136.7, 136.2, 133.9, 131.9,

130.9, 129.5, 127.9, 120.4, 68.1, 64.4, 63.2, 53.4, 36.1, 33.7; MS (ESI+) for

$\text{C}_{22}\text{H}_{19}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 524.0 ($\text{M}+\text{H}$) $^+$. MS (ESI-) for $\text{C}_{22}\text{H}_{19}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 522.0 ($\text{M}-\text{H}$) $^-$.

Anal. Calcd for $\text{C}_{22}\text{H}_{19}\text{Cl}_2\text{N}_3\text{O}_6\text{S} \cdot 0.31 \text{ H}_2\text{O}$: C, 49.86; H, 3.73; N, 7.93. Found: C, 49.61;

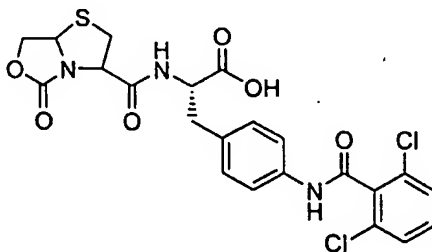
10 H, 3.82; N, 7.54. % Water (KF): 1.06.

Example 282.

4-[(2,6-Dichlorobenzoyl)amino]-*N*-[(tetrahydro-5-oxo-5*H*-thiazolo[3,2-*c*]oxazol-3-yl)-

carbonyl]-L-phenylalanine (more polar diastereomer)

15 (Scheme Q, Q-8: where $\text{R}_{\text{Q},1}$ and $\text{R}_{\text{Q},2}$ are equal to hydrogen, Z is O, R_5 is 4-[(2,6-dichlorobenzoyl)amino]phenyl, and stereochemistry of the C-terminal amino acid is (*S*)).



20 Example 282 (diastereomer of Example 281) was prepared as described in Scheme

Q. Physical data as follows: IR (drift) 3296, 1753, 1666, 1603, 1579, 1535, 1517, 1431,

1411, 1390, 1324, 1267, 1207, 798, 781 cm^{-1} ; ^1H NMR (300 MHz, $\text{DMSO}-d_6$) δ 8.57 (1

H), 7.54 (5 H), 7.20 (2 H), 5.23 (1 H), 4.86 (1 H), 4.66 (1 H), 4.45 (2 H), 3.22 (1 H), 3.09

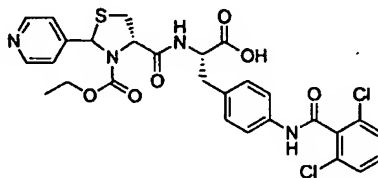
(1 H), 2.93 (m, 2 H); ^{13}C NMR (75 MHz, CD_3OD) δ 172.8, 169.7, 163.8, 160.6, 136.8,

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136.2, 133.7, 131.9, 130.9, 129.5, 127.9, 120.2, 68.1, 64.9, 63.1, 53.6, 37.5, 36.2, 34.2;
 MS (ESI+) for $C_{22}H_{19}Cl_2N_3O_6S$ m/z 523.9 (M+H)⁺. MS (ESI-) for $C_{22}H_{19}Cl_2N_3O_6S$ m/z
 521.9 (M-H)⁻; HRMS (EI) calcd for $C_{22}H_{19}Cl_2N_3O_6S$ 523.0372, found 523.0366. Anal.
 Calcd for $C_{22}H_{19}Cl_2N_3O_6S \cdot 0.35 H_2O$: C, 49.79; H, 3.74; N, 7.92. Found: C, 50.14; H,
 5 4.08; N, 8.13. % Water (KF): 1.19.

Example 283.

10 (4S)-4-[[[(1S)-1-Carboxy-2-[4-[(2,6-
 dichlorobenzoyl)amino]phenyl]ethyl]amino]carbonyl]-2-(4-pyridinyl)-3-
 thiazolidinecarboxylic acid 3-ethyl ester



15

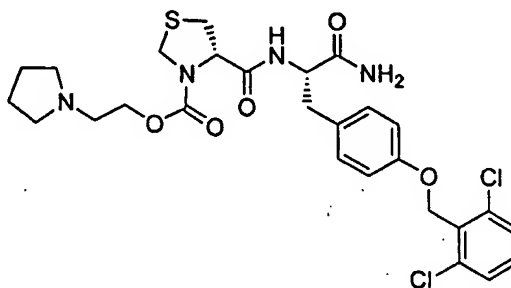
Example 283 was prepared as described in Scheme B using D-cysteine and 4-
 pyridinecarboxaldehyde to form the requisite thiazolidine carboxylic acid. Physical data
 as follows: IR (drift) 3055, 2981, 2928, 1679, 1604, 1535, 1515, 1450, 1431, 1406, 1378,
 1331, 1194, 797, 778 cm^{-1} ; ¹H NMR (300 MHz, CD₃OD) δ 8.68 (2 H), 8.28 (2 H), 7.55 (2
 20 H), 7.42 (4 H), 7.25 (2 H), 6.28 (1 H), 4.70 (1 H), 4.09 (2 H), 3.46 (1 H), 3.16 (1 H), 2.94
 (1 H), 2.73 (1 H), 1.17 (3 H); ¹³C NMR (75 MHz, CD₃OD) δ 190.2, 171.1, 162.1, 153.8,
 141.4, 135.3, 134.6, 131.9, 130.3, 129.4, 128.2, 126.4, 122.4, 118.5, 61.0, 51.7, 35.5, 26.6,
 11.7; MS (FAB) m/z (rel. intensity) 617 (MH⁺, 99), 621 (35), 620 (72), 619 (99), 618
 (91), 617 (99), 371 (22), 179 (23), 173 (28), 124 (27), 57 (24); HRMS (FAB) calcd for
 25 $C_{28}H_{26}Cl_2N_4O_6S + H$, 617.1028, found 617.1019. Anal. Calcd for $C_{28}H_{26}Cl_2N_4O_6S \cdot 0.9$
 HCl $\cdot 1.1 H_2O$: C, 50.19; H, 4.38; N, 8.36; Cl, 15.34. Found: C, 49.79; H, 4.49; N, 8.11;
 Cl, 15.05. % Water (KF): 2.96.

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Example 284.

4-[[[(1*S*)-2-Amino-1-[4-[(2,6-dichlorophenyl)methoxy]phenyl]methyl]-2-oxoethyl]amino]-carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(1-pyrrolidiny)ethyl] ester

5



Example 284 was prepared as described in Scheme C. Physical data as follows:

IR (drift) 1709, 1675, 1511, 1458, 1435, 1421, 1390, 1380, 1354, 1299, 1240, 119

5, 1179, 1114, 765 cm^{-1} ; ^1H NMR (300 MHz, CD_3OD) δ 8.40 (1 H), 7.41 (3 H), 7.22 (2

10 H), 6.99 (2 H), 5.29 (2 H), 4.46 (6 H), 3.77 (2 H), 3.50 (2 H), 3.19 (4 H), 2.89 (1 H), 2.04 (1 H), 2.12 (4 H); ^{13}C NMR (75 MHz, CD_3OD) δ 174.5, 171.6, 171.1, 157.9, 153.3, 136.6,

132.2, 130.6, 130.0, 129.7, 128.3, 114.5, 64.9, 63.2, 62.1, 60.9, 54.4, 53.6, 50.0, 49.0,

37.0, 36.5, 35.2, 22.6; MS (ESI+) for $\text{C}_{27}\text{H}_{32}\text{Cl}_2\text{N}_4\text{O}_5\text{S}$ m/z 595.1 (M+H) $^+$. HRMS (FAB)

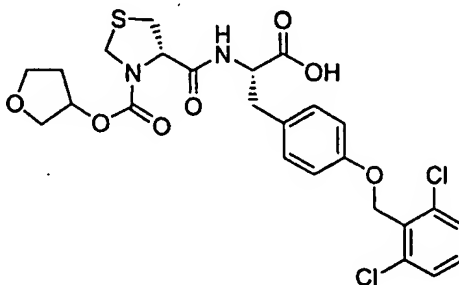
calcd for $\text{C}_{27}\text{H}_{32}\text{Cl}_2\text{N}_4\text{O}_5\text{S} + \text{H}$, 595.1548, found 595.1531.

15

Example 285.

(4*S*)-4-[[[(1*S*)-1-Carboxy-2-[4-[(2,6-dichlorophenyl)methoxy]phenyl]ethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid 3-(3-tetrahydrofuranyl) ester

20



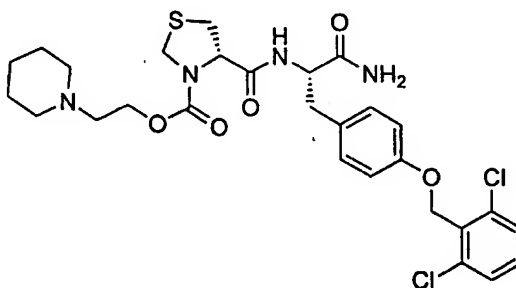
-242-

Example 285 was prepared as described in Scheme A. Physical data as follows:

¹H NMR (300 MHz, CD₃OD) δ 7.41 (3 H), 7.18 (2 H), 6.98 (2 H), 5.28 (2 H), 5.15 (1 H)
4.69 (2 H), 4.44 (1 H), 3.82 (4 H); ¹³C NMR (75 MHz, DMSO-*d*₆) δ 172.8, 157.9, 136.7,
132.2, 130.6, 130.1, 129.5, 128.3, 114.5, 76.8, 72.7, 66.5, 64.9, 62.4, 53.5, 36.3, 34.9,
32.4; MS (FAB) *m/z* (rel. intensity) 569 (MH⁺, 75), 571 (51), 570 (29), 569 (75), 322
(22), 161 (20), 159 (30), 89 (25), 73 (31), 71 (99), 43 (41); HRMS (FAB) calcd for
C₂₅H₂₆Cl₂N₂O₇S + H₁ 569.0916, found 569.0939. Anal. Calcd for C₂₅H₂₆Cl₂N₂O₇S: C,
52.73; H, 4.60; N, 4.92;. Found: C, 52.41; H, 4.80; N, 4.62.

Example 286.

(4*S*)-4-[[[(1*S*)-2-Amino-1-[[4-[(2,6-dichlorophenyl)methoxy]phenyl]methyl]-2-oxoethyl]-
amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(1-piperidiny)ethyl] ester



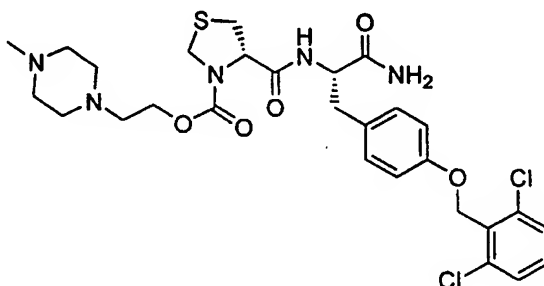
Example 286 was prepared as described in Scheme C. Physical data as follows:

¹H NMR (300 MHz, CD₃OD) δ 7.43 (3 H), 7.22 (2 H), 6.98 (2 H), 5.28 (2 H), 4.66 (2 H),
4.59 (1 H), 4.47 (1 H), 4.47 (2 H), 3.05 (10 H), 1.78 (4 H), 1.62 (2 H); ¹³C NMR (75 MHz,
CD₃OD) δ 175.9, 172.9, 159.3, 138.1, 133.6, 132.0, 131.4, 131.1, 129.7, 120.1, 115.9,
66.3, 64.6, 63.7, 62.3, 61.9, 57.8, 55.8, 55.1, 51.3, 50.3, 38.5, 37.8, 36.6, 25.0, 23.5; MS
(ESI⁺) for C₂₈H₃₄Cl₂N₄·O₅S *m/z* 609.0 (M+H)⁺.

Example 287.

(4*S*)-4-[[[(1*S*)-2-Amino-1-[[4-[(2,6-dichlorophenyl)methoxy]phenyl]methyl]-2-oxoethyl]-amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(4-methyl-1-piperazinyl)ethyl] ester

5



Example 287 was prepared as described in Scheme C. Physical data as follows:

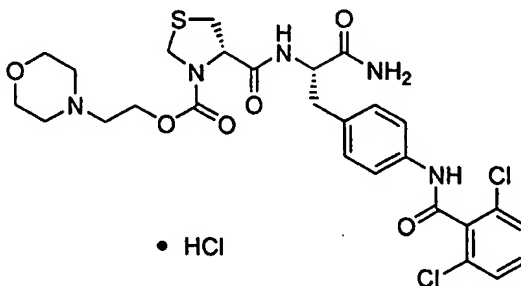
¹H NMR (300 MHz, CD₃OD) δ 7.41 (3 H), 7.21 (2 H), 6.98 (2 H), 5.28 (2 H), 4.67 (1 H), 4.58 (1 H), 4.46 (1 H), 4.23 (2 H), 3.20 (2 H), 2.71 (13 H), 2.50 (3 H); MS (ESI+) for C₂₈H₃₅Cl₂N₅O₅S *m/z* 624.0 (M+H)⁺; Anal. Calcd for C₂₈H₃₅Cl₂N₅O₅S • 0.5 C₂H₄O₂ • 0.5 H₂O: C, 49.82; H, 5.54; N, 10.02. Found: C, 49.82; H, 5.77; N, 9.65.

10

Example 288.

(4*S*)-4-[[[(1*S*)-2-Amino-1-[[4-[(2,6-dichlorobenzoyl)amino]phenyl]methyl]-2-oxoethyl]-amino]carbonyl]-3-thiazolidinecarboxylic acid 3-[2-(4-morpholinyl)ethyl] ester

15

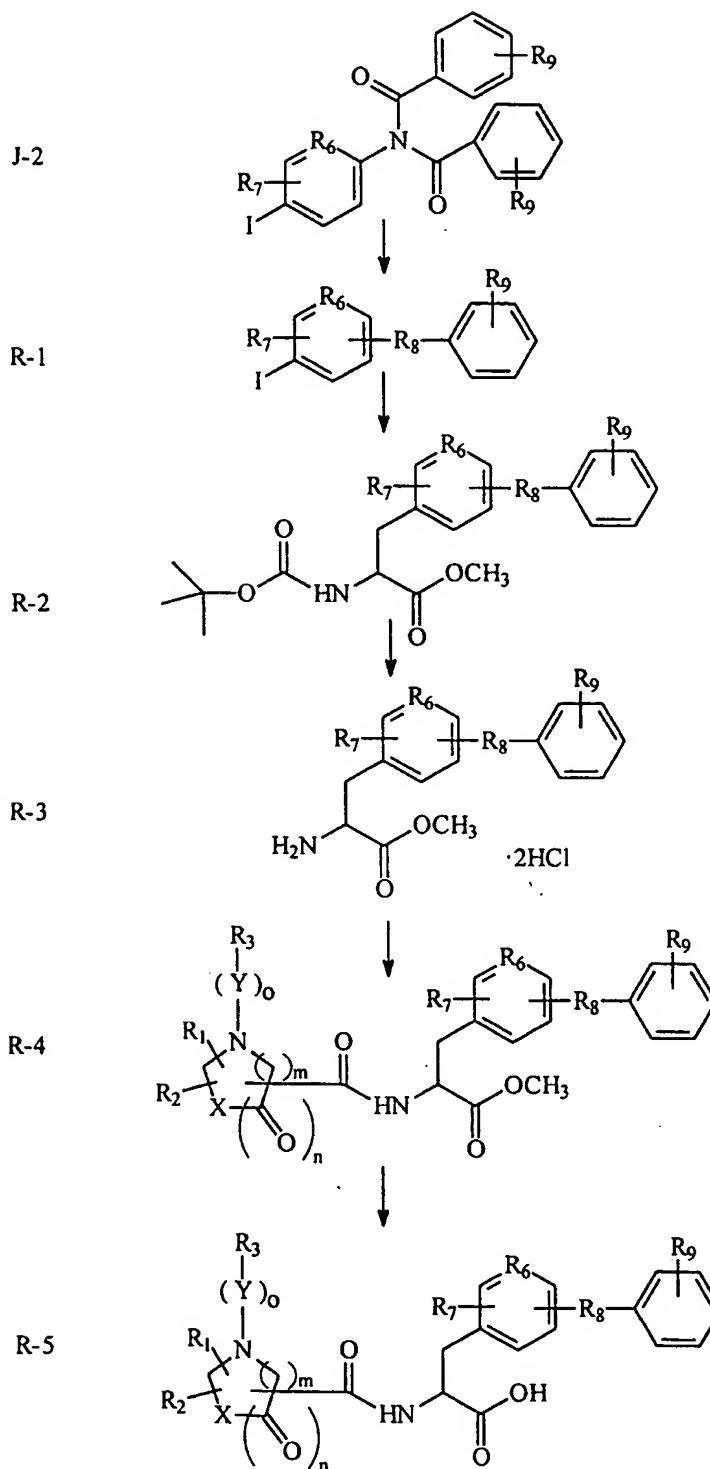


Example 288 was prepared as described in Scheme C. Physical data as follows:

IR (drift) 1671, 1603, 1536, 1518, 1430, 1415, 1361, 1349, 1324, 1269, 1194, 1134, 1118, 1104, 799 cm^{-1} ; ^1H NMR (300 MHz, CD_3OD) δ 7.62 (1 H), 7.46 (3 H), 7.31 (2 H), 4.68 (3 H), 4.41 (3 H), 3.95 (4 H), 3.48 (5 H), 3.22 (3 H), 2.90 (2 H); ^{13}C NMR (75 MHz, CD_3OD)
5 δ 175.8, 173.1, 165.3, 154.7, 138.2, 137.6, 135.2, 133.3, 132.4, 130.9, 129.4, 121.7, 121.5, 65.0, 64.8, 63.7, 60.6, 57.4, 55.8, 53.6, 51.4, 50.4, 38.6, 38.1, 36.6, 35.0; MS (ESI+) for $\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}$ m/z 623.9 ($\text{M}+\text{H}$) $^+$; HRMS (FAB) calcd for $\text{C}_{27}\text{H}_{31}\text{Cl}_2\text{N}_3\text{O}_6\text{S}+\text{H}$ 624.1450, found 624.1452.

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Scheme R.



Scheme R

Scheme R teaches a general method for the preparation of Examples corresponding to structures R-4 and R-5, where R_6 is nitrogen. Thus reaction of the amide R-1 (obtained from the imide J-2), with the organozinc derived from a suitable protected β -iodoalanine, provides the acylamino azaphenylalanine R-2. *N*-deprotection of R-2 gives the aminoester R-3, that is used (as exemplified by reagent A-4 of Scheme A, and by reagent B-5 of Scheme B) for the synthesis of Examples R-4 and R-5 of this invention.

10

Preparation 87

(Scheme R: R-1 where R_6 is N, R_7 is H, R_8 is -NHC(O)-, and R_9 is 2,6-dichloro) **2,6-Dichloro-*N*-(5-iodo-2-pyridinyl)benzamide** ($C_{12}H_7Cl_2IN_2O$). A mixture of J-2 and $NH_2NH_2 \cdot H_2O$ in MeOH is refluxed for 6 h under Ar. The reaction mixture is cooled, and the MeOH is removed *in vacuo*. The residue is partitioned between H_2O and EtOAc. The EtOAc extracts are dried, filtered and concentrated to give a brown-colored solid, that is purified by silica flash chromatography (99:1 toluene/EtOAc) to provide **Preparation 87**: TLC (98:2 toluene/EtOAc) R_f = 0.43; 1H NMR ($CDCl_3$, 300 MHz) δ 10.23 (1H), 8.26 (1H), 7.99 (1H), 7.57 (1H), 7.41–7.31 (3H); ^{13}C NMR ($CDCl_3$, 75 MHz) δ 163.40, 153.22, 150.95, 147.16, 135.83, 132.72, 131.90, 130.34, 128.65, 128.50, 117.09, 86.55; MS (ESI) 393, 391.

20

Preparation 88

(Scheme R: R-2 where R_6 is N, R_7 is H, R_8 is -NHC(O)-, R_9 is 2,6-dichloro and the stereochemistry is *S*)

(*S*)-6-[(2,6-Dichlorobenzoyl)amino]- α -[[[(1,1-dimethylethoxy)carbonyl]amino]-3-pyridinepropanoic acid methyl ester ($C_{21}H_{23}Cl_2N_3O_5$). To an amberized flask containing activated Zn dust (0.0802 g, 12.27 mmol) under Ar is added dry THF (6 mL) and 1,2-dibromoethane (0.045 mL). This suspension is brought briefly to a gentle reflux, and then is cooled to rt. A solution of TMSCl (1 M in THF, 0.39 mL) is added. The reaction mixture is stirred at 45 ± 5 °C for 30 min, and then is cooled to rt. To this mixture is added

30

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a degassed solution of *N*-[(1,1-dimethylethoxy)carbonyl]-3-iodo-L-alanine methyl ester (4.04 g, 12.27 mmol) in 2:1 *N,N*-dimethylacetamide/THF (18 mL). The reaction mixture is stirred at 45 ± 5 °C for 5 h, and then is cooled to 0 °C. To this mixture is added $\text{PdCl}_2(\text{PPh}_3)_2$ (0.428 g) followed immediately by a degassed solution of **Preparation 87** in 1:1 *N,N*-dimethylacetamide/THF (19 mL). This reaction mixture is stirred at 45 ± 5 °C for 44 h. It is cooled to 0 °C, and is quenched with cold aqueous satd NH_4Cl . This mixture is extracted with EtOAc. The combined EtOAc extracts are washed with aqueous satd NH_4Cl and brine. The EtOAc extracts are dried, filtered and concentrated to provide a green-brown colored oil, that is purified by silica flash chromatography (steps of 750:250:1, 700:300:1 and 650:350:1 heptane/EtOAc/*i*PrOH) to provide **Preparation 88**: TLC R_f = 0.28 (7:3 hexanes/EtOAc).

Preparation 89

(Scheme R: R₆ is N, R₇ is H, R₈ is -NHC(O)-, R₉ is 2,6-dichloro and the stereochemistry is *S*)

(*S*)- α -Amino-6-[(2,6-dichlorobenzoyl)amino]-3-pyridinepropanoic acid methyl ester dihydrochloride salt (C₁₆H₁₅Cl₂N₃O₃·2HCl, R-3). A solution of **Preparation 88** (0.812 g, 1.73 mmol) in 4 M HCl in dioxane (20 mL) is stirred under Ar at rt for 20 h. The reaction mixture is concentrated *in vacuo*, and the residue is taken up in H₂O (60 mL). This aqueous mixture is extracted with Et₂O (3 60 mL), and the Et₂O extracts are discarded. The aqueous solution is frozen and lyophilized to provide **Preparation 89**: IR (diffuse reflectance) 3021, 2995, 2953, 2893, 2884, 2866, 2853, 2844, 2341, 2015, 1916, 1749, 1646, 1569, 1252 cm⁻¹; MS (EI) 367 (M⁺), 282, 280, 262, 175, 173, 147, 145, 109, 107, 88.

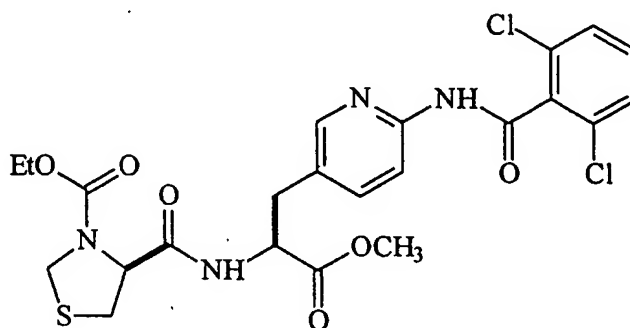
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Preparation 90 & Example 289

[*S*-(*R**,*R**)]-4-[[[1-[[2-[(2,6-Dichlorobenzoyl)amino]-5-pyridinyl]methyl]-2-methoxy-2-oxoethyl]amino]carbonyl]-3-thiazolidinecarboxylic acid ethyl ester
(C₂₃H₂₄Cl₂N₄O₆S)

(Scheme R: R₁ is H, R₂ is H, R₃ is -Et, X is S, (Y) is -C(O)O, m is 2, n is 0, o is 1, R₆ is N, R₇ is H, R₈ is -NHC(O)-, R₉ is 2,6-dichloro and the stereochemistry is [*S*-(*R**,*R**)])



To a mixture of the *N*-acylthiazolidinecarboxylic acid (0.292 g, 1.42 mmol), HOAt (0.193 g, 1.42 mmol) in 4:1 CH₂Cl₂/DMF (5.25 mL) at 0 °C is added EDC (0.272 g, 1.42 mmol). This reaction mixture is stirred at 0 °C for 20 min. Solid **Preparation 89** (0.568 g, 1.29 mmol) and NMM (0.316 mL, 3.27 mmol) are added. The resulting reaction mixture is stirred at 0 °C for 4 h, and then is kept at 4 °C for 40 h. The mixture is concentrated *in vacuo*, and the residue is taken up in CH₂Cl₂. The CH₂Cl₂ mixture is extracted with H₂O, aq satd NaHCO₃, and H₂O. The combined aqueous washes are back-extracted with CH₂Cl₂. The combined CH₂Cl₂ extracts are dried, filtered and concentrated to a pale yellow-colored foam, that is purified by silica flash chromatography (600:400:1 EtOAc/heptane/iPrOH) to give **Preparation 90 (Example 289)**: IR (diffuse reflectance) 3275, 1742, 1697, 1665, 1586, 1559, 1532, 1476, 1427, 1398, 1382, 1351, 1344, 1311,

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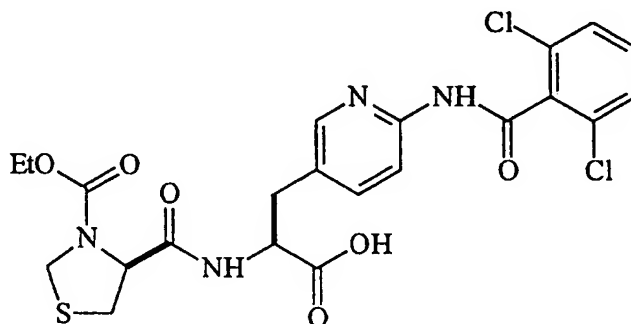
1284, 1279, 1242, 1223, 1196, 1100, 1023, 802, 787, 770, 697 cm^{-1} ; MS (EI) 554 (M^+), 495, 422, 395, 352, 315, 293, 280, 172, 160, 144, 116, 107, 88, 60.

Example 290

5 $[S-(R^*, R^*)]-4-[[[1\text{-Carboxy-2-[2-[(2,6-dichlorobenzoyl)amino]-5-pyridinyl]ethyl]amino]carbonyl]-3\text{-thiazolidinecarboxylic acid ethyl ester}$
 $(\text{C}_{22}\text{H}_{22}\text{Cl}_2\text{N}_4\text{O}_6\text{S}, \text{Example 290})$

(Scheme R: R-4 where R_1 is H, R_2 is H, R_3 is -Et, X is S, (Y) is -C(O)O, m is 2, n is 0, o is 1, R_6 is N, R_7 is H, R_8 is -NHC(O)-, R_9 is 2,6-dichloro and the stereochemistry is [$S-(R^*, R^*)$])

10



To a solution of **Preparation 289** (0.400 g, 0.72 mmol) in 6:1 THF/ H_2O (25.6 mL) at 0 °C under Ar is added slowly over 4 h (via syringe pump) an aqueous solution of NaOH (1 M, 7.92 mL). The reaction mixture is stirred an additional 1.5 h. The reaction mixture is partitioned between aqueous HCl and EtOAc. The aqueous solution is separated, and is extracted further with EtOAc. The combined EtOAc extracts are dried, filtered and concentrated to a beige-colored foam. This foam is taken up in 1:1 MeCN/ H_2O . This solution is frozen and lyophilized to give **Example 290** as a beige-colored solid: mp 142–144 °C; IR (diffuse reflectance) 3169, 3094, 3031, 2980, 2964, 2935, 1735, 1691, 1591, 1556, 1531, 1480, 1431, 1400, 1379, 1344, 1308, 1288, 1266, 1216, 1194, 1148, 799, 782, 772 cm^{-1} ; Anal. C 48.72, H 4.29, Cl 12.26, N 9.95, S 5.62 (calcd C 48.81, H 4.10, Cl 13.10, N 10.35, S 5.92).

15

20

Biological Assays

Jurkat-Endothelial Cell Adhesion Assay:

The following assay established the activity of the present compounds in inhibiting β_1 -mediated cell adhesion in a representative *in vitro* system. This assay
5 measures the adhesive interactions of a T-cell line, Jurkat, known to express the $\alpha_4\beta_1$ integrin, to endothelial monolayers in the presence of test compounds. The test compounds were added in increasing concentrations to T-cells and then the T-cell compound mixture was added to IL-1 stimulated endothelial cell monolayers. The plates were incubated, washed and the percentage of attached cells was
10 quantitated. The present assay directly demonstrates the cell adhesion inhibitory activity and adhesion modulatory activity of the compounds.

Human umbilical vein endothelial cells were purchased from Clonetics (San Diego, CA.) at passage number 2. The cells were grown on 0.5% porcine skin gelatin pre-coated flasks (Sigma, St. Louis MO.) in EGM-UV media
15 (Clonetics, San Diego, CA) supplemented with 10% fetal bovine serum. Cells are refed every 2-3 days reaching confluence by day 4 to 6. The cells are monitored for factor VIII antigen and results show that at passage 12, the cells are positive for this antigen. The endothelial cells are not used following passage 6.

The T-cell line Jurkat was obtained from American Type Tissue Culture
20 Collection (Rockville, MD) and the cells were cultured in RPMI containing 10 % fetal calf serum. The cells were washed twice in Hank's Balanced Salt Solution (HBSS) and resuspended in Dulbecco's Minimal Eagle's Media (DMEM) containing 2.5 mg/ml Human Serum Albumin (HSA). Jurkat cells (1×10^6 cells/ml) were stained with 10 ng/ml BCECF-AM (Molecular Probes, Eugene, OR)) in
25 HBSS without phenol red. The cells were loaded with BCECF for 60 minutes in the dark at 37°C, washed 2 times, and resuspended in DMEM-HSA solution.

Confluent endothelial monolayers, grown in 96-well tissue culture plates, were stimulated for 4 hr. at 37 °C with 0.1 ng/ml (~50 U/ml) recombinant IL-1 (Amgen, Thousand Oaks, CA). Following this incubation, the monolayers were
30 washed twice with HBSS and 0.1 ml of DMEM-HSA solution was added. Jurkat

cells (5×10^5 cells) were combined with the appropriate concentration of the test compound and 0.1 ml of the Jurkat cell-compound mixture was added to the endothelial cell monolayers. Generally, 100, 20, 5 and 1.25 μ M compound concentrations were tested. These concentrations are adjusted downward for
5 analogs found or thought to be more potent. The plates were placed on ice for 5 minutes to allow for Jurkat cell settling and the plates were incubated at 37 °C for 20 minutes. Following this incubation, the monolayers were washed twice with PBS containing 1 mM calcium chloride and 1 mM magnesium chloride and the plates were read using a Millipore Cytofluor 2300 (Marlboro, MA.). Fluorescence
10 in each well was measured as Arbitrary Fluorescence Units and percent adhesion in the absence of compound was adjusted to 100% and the % adhesion in the presence of compound was calculated. Monolayers were also fixed in 3% paraformaldehyde and evaluated microscopically to verify the adhesion. This procedure is a modification of a previously published method (Cardarelli et al., *J. Biol. Chem.* 269:18668-18673 (1994)).
15

Jurkat-CS-1 assay

The CS-1 derived peptide, CLHPGEILDVPST, and the scrambled control peptide, CLHGPIELVSDPT, were synthesized on a Beckman 990 synthesizer
20 using t-Boc methodology. The peptides were immobilized onto microtiter plates using the heterobifunctional crosslinker 3-(2-pyridyldithio)propionic acid N-hydroxysuccinimide ester (SPDP) as reported by Pierschbacher et al., *Proc. Natl. Acad. USA*, 80:1224-1227 (1983). Microtiter plates were coated with 20 μ g/ml HSA for 2 hr. at room temperature, washed once with PBS and derivatized with 10
25 μ g/ml SPDP for 1 hr. After washing, 100 μ l of a 100 μ g/ml cysteine containing peptide solution which had been recently dissolved was added to the wells and allowed to crosslink to the plates overnight at 4 °C. Unbound peptide was removed from plates by washing with PBS. To block non-reacted sites, the plates are coated with 100 μ l of a 2.5 mg/ml BSA solution in PBS for 1 hr. at 37 °C. 100 μ l of
30 Jurkat cells (2.5×10^6 cells/ml) in DMEM plus BSA (2.5 mg/ml) was mixed with

an appropriate concentration of the compound to be tested and the mixture was added to peptide coated dishes and incubated for 1 hr. at 37 °C. Generally 100, 20, 5 and 1.25 μ M concentrations of the compound were tested. The concentrations of the compound were adjusted downward for compounds thought or found to be more potent.

Following this incubation the plates were washed once with PBS and the attached cells were fixed with 3% paraformaldehyde in PBS and stained with 0.5% toluidine blue in 3.7% formaldehyde. The cells were stained overnight at room temperature and the optical density at 590 nm of toluidine blue stained cells was determined using a vertical pathway spectrophotometer to quantitate attachment (VMAX Kinetic Microplate Reader, Molecular Devices, Menlo Park, CA). This procedure is a modification of a previously published method (Cardarelli et al, J. Biol. Chem., 269:18668-18673 (1994) and Cardarelli et al, Proc. Natl. Acad. Sci. USA, 83:2647-2651 (1986)).

The preferred compounds are those which have low IC_{50} values in the Jurkat EC assay or the Jurkat-CS-1 assay described above or which have at least moderate activity in both assays. All of the compounds of the present invention have an activity of less than 50 μ M in the Jurkat CS-1 assay or less than 500 μ M in the Jurkat EC assay. Compounds with activity in the Jurkat CS-1 assay preferably have IC_{50} values of less than 1 μ M, more preferably less than 0.5 μ M, most preferably less than or equal to 0.08 μ M. Compounds with activity in the Jurkat EC assay preferably have IC_{50} values of less than 10 μ M, more preferably less than 5 μ M, most preferably less than or equal to 0.8 μ M.

In the Jurkat EC Assay, IC_{50} value ranges (μ M) are depicted by A, B, and C and in the Jurkat CS-1 Assay, IC_{50} value ranges are depicted by D, E, and F. These ranges are as follows:

In vitro data:

EC: $A \geq 1 \mu M$; $1 \mu M > B > 0.25 \mu M$; $C \leq 0.25 \mu M$

CS-1 $D \geq 0.75 \mu M$; $0.75 \mu M > E > 0.05 \mu M$; $F \leq 0.05 \mu M$.

Ex.	JK/EC	JK/CS-1
1	A	E
2	C	E
3	C	F
4	C	F
5	B	E
6	A	D
7	A	E
8	B	E
9	C	F
10	C	F
11	A	E
12	C	F
13	A	D
14	B	E
15	A	D
16	C	E
17	A	D
18	A	D
19	C	E
20	A	D
21	C	E
22	A	D
23	C	F
24	B	E
25	C	F
26	A	D
27	A	D
28	A	D
29	B	E
30	A	E
31	C	F
32	A	D
33	C	E
34	A	E
35	B	E
36	C	F
37	C	F
38	A	D
39	C	E
41	A	D
42	B	E

Ex.	JK/EC	JK/CS-1
43	C	F
44	A	D
45	B	E
46	A	D
47	C	F
48	A	D
49	A	D
50	A	E
51	B	E
52	C	E
53	C	F
54	A	E
55	C	F
56	A	E
57	A	D
58	C	F
59	A	E
60	C	F
61	B	E
62	C	F
63	C	F
64	C	F
65	A	E
66	C	E
67	A	E
68	A	E
69	A	D
70	B	F
71	A	D
72	A	D
73	A	E
74	B	E
75	A	D
76	C	E
77	A	D
78	A	E
79	A	D
80	A	D
82	B	E
83	A	E
84	C	F

Ex.	JK/EC	JK/CS-1
85	C	F
86	A	D
87	A	D
88	A	E
89	A	E
90	A	
91	A	D
92	A	D
93	A	D
94	A	
95	A	
96	A	D
97	A	D
99	C	F
100	B	E
101	B	E
102	A	D
103	B	F
104	A	E
105	A	E
107	A	E
109	A	D
110	A	
111	B	E
112	A	
113	A	E
114	A	D
115	A	E
116	A	D
117	A	E
119	A	D
121	A	E
123	A	D
124	A	D
125	B	E
126	A	D
127	C	E
129	B	E
131	B	E
133	A	D
135	B	E

Ex.	JK/EC	JK/CS-1
137	B	E
139	A	E
140	A	D
141	A	E
142	A	D
143	A	E
144	A	E
145	A	D
146	A	D
147	A	D
148	C	F
149	C	F
150	B	E
151	C	F
152	C	F
153	A	D
154	B	F
155	A	D
156	A	E
157	C	E
158	A	D
159	A	D
161	A	D
163	A	
164	A	
165	C	
166	A	
167	B	E
168	A	
169	A	
170	A	
171	A	E
172	C	F
173	C	E
174	B	
175	A	
176	A	E
177	A	F
178	A	F
179	B	
180	A	

Ex.	JK/EC	JK/CS-1
181	A	E
182	B	E
183	A	
184	C	F
185	C	F
186	C	F
187	A	E
188	A	E
189	A	E
190	B	E
191	A	E
192	C	F
193	C	E
195	A	E
196	A	E
197	A	D
298	B	
200	A	E
201	A	E
202	A	
203	A	E
204	B	E
205	B	
206	B	E
207	A	E
208	A	
209	A	
210	A	
211	A	
212	A	
213	A	
214	A	
215	A	
216	A	E
217	A	
218	A	
219	A	
220	A	E
221	A	
222	A	E
224	A	E

Ex.	JK/EC	JK/CS-1
225	A	E
226	A	E
227	A	E
228	A	E
229	A	E
230	A	E
231	A	E
232	A	D
233	A	E
234	A	E
235	A	E
236	A	D
237	A	E
238	A	E
239	A	E
240	A	E
241	A	E
242	A	E
243	B	E
244	B	E
245	C	E
246	B	E
247	B	E
248	B	E
249	B	E
250	C	E
251	C	F
252	C	F
253	C	E
254	B	E
255	A	E
256	C	F
257	B	E
258	C	F
259	C	E
260	B	E
261	B	E
262	C	F
263	B	E
264	C	F
265	C	F

Ex.	JK/EC	JK/CS-1
266	C	F
267	A	E
268		D
269		D
270		E
271		E
272		E
273		D
274		E
275		F
276		D
277		D
278		E
279		E
280		E
281		E
282		F
283		E
284		E
285		E
286		E
287		E
288		D
290		F

Biological Activity in Dextran Pleurisy Model

Certain compounds of the present invention were tested in a Dextran® pleurisy model.

5 Rationale for Developing an $\alpha_4\beta_1$ Integrin Antagonist to Treat Inflammatory Diseases

VLA-4, a member of the β_1 integrin family of adhesion molecules, is thought to play a critical role in several types of inflammatory disease processes by promoting leukocyte adhesion to vascular cell adhesion molecule (VCAM-1) and the CS-1 domain of fibronectin in extracellular tissue matrix (Elices MJ, Osborn L, Takada Y, Crouse C,

- 10 Luhowskyj S, Hemler M, Lobb RR. VCAM-1 on activated endothelium interacts with the leukocyte integrin VLA-4 at a site distinct from the VLA-4-fibronectin binding site. Cell; 60: 577-584. 1990. Humphries MJ, Akiyama SK, Komoriya A, Olden K, Yamada KM.

- Identification of an alternatively-spliced site in human plasma fibronectin that mediates cell type-specific adhesion. J Cell Biol; 103: 2637-2647, 1986, Wayner EA, Garcia-Pardo A, Humphries MJ, McDonald JA, Carter WG. Identification and characterization of the T lymphocyte adhesion receptor for an alternative cell attachment domain (CS-1) in plasma
- 5 fibronectin. J Cell Biol; 109: 1321-1330, 1989, Guan J-L, Hynes RO. Lymphoid cells recognize an alternatively-spliced segment of fibronectin via the integrin $\alpha_4\beta_1$. Cell; 60: 53-61, 1990) Of the cell types expressing VLA-4, the major emphasis has been on eosinophils, lymphocytes, and monocytes. Validation of the role of VLA-4 has relied predominantly on the use of anti-VLA-4 antibodies which have been shown to suppress
- 10 delayed-type hypersensitivity responses (Issekutz TB. Dual inhibition of VLA-4 and LFA-1 maximally inhibits cutaneous delayed-type hypersensitivity-induced inflammation. Am J Pathol; 143: 1286-1293, 1993, Scheynius A, Camp RL, Puré E. Reduced contact sensitivity reactions in mice treated with monoclonal antibodies to leukocyte function-associated molecule-1 and intercellular adhesion molecule-1. J Immunol; 150: 655-663,
- 15 1993, Ferguson TA, Kupper TS. Antigen-independent processes in antigen-specific immunity. J Immunol; 150: 1172-1182, 1993, Chisholm PL, Williams CA, Lobb RR. Monoclonal antibodies to the integrin α_4 subunit inhibit the murine contact hypersensitivity response. Eur J Immunol; 23: 682-688, 1993. Elices MJ, Tamraz S, Tollefson V, Vollger LW. The integrin VLA-4 mediates leukocyte recruitment to skin
- 20 inflammatory sites *in vivo*. Clin Exp Rheumatol; 11 (Suppl 8) S77-80, 1993, experimental allergic encephalomyelitis (Yednock TA, Cannon C, Fritz LC, Sanchez-Madrid F, Steinman LM, Karin N. Prevention of experimental autoimmune encephalomyelitis by antibodies against $\alpha_4\beta_1$ integrin. Nature; 356: 63-66, 1992, Canella B, Raine CS. The VCAM-1/VLA-4 pathway is involved in chronic lesion expression in
- 25 multiple sclerosis (MS). J Neuropathol Exp Neurol; 52: 311, 1993), HIV-induced encephalitis (Sasseville VG, Newman W, Brodie SJ, Hesterberg P, Pauley D, Ringler DJ. Monocyte adhesion to endothelium in simian immunodeficiency virus-induced AIDS encephalitis is mediated by vascular cell adhesion molecule-1/ $\alpha_4\beta_1$ integrin reactions. Am J Pathol; 144: 27-40, 1994), pulmonary inflammation and airway hyperactivity in asthma
- 30 (Abraham WM, Sielczak MW, Ahmed A, Cortes A, Lauredo IT, Kim J, Pepinsky, B, et al.

α -integrins mediate antigen-induced late bronchial responses and prolonged airway hyperresponsiveness in sheep. J Clin Invest; 93: 776-787, 1994. Pretolani M, Ruffié C, Roberto LapaeSilva J, Joseph D, Lobb RR, Vargaftig BB. Antibody to very late activation antigen 4 prevents antigen-induced bronchial hyperreactivity and cellular infiltration in the guinea-pig airways. J Exp Med; 180: 795-805, 1994), experimental models of autoimmune-mediated diabetes (Yang X-D, Karin N, Tisch R, Steinman L, McDevitt HO. Inhibition of insulinitis and prevention of diabetes in non-obese diabetic mice by blocking L-selectin and very late antigen 4 adhesion receptors. Proc Natl Acad Sci USA; 90: 10494-10498, 1993, Burkly LC, Jakubowski A, Hattori M. Protection against adoptive transfer of autoimmune diabetes mediated through very late antigen-4 integrin. Diabetes; 43: 529-534, 1994), and experimental colitis (Podolsky DK, Lobb R, King N, Benjamin CD, Pepinsky B, Sehgal P, et al. Attenuation of colitis in the cotton-top Tamarin by anti-4 integrin monoclonal antibody. J Clin Invest; 92: 372-380, 1993). Since eosinophils represent a major component of the inflammatory cell influx in asthmatic lung tissue we developed a simple acute inflammatory model of VLA-4 integrin-dependent eosinophil infiltration which could be used to identify VLA-4 antagonists; such compounds would be of potential value in the treatment of asthma as well as other diseases in which VLA-4 played a role.

Materials and Methods

20 Animals, housing and viral testing:

C57BL/6 mice (Jackson, Bar Harbor, ME;), 6-8 weeks old, weighing 20-25g were used throughout. All mice were acclimated for at least 7-14 days after arrival and maintained under controlled temperature (20-22°C) and a 12 hr daily light cycle (6.00 A.M. - 6.00 P.M.). Mice were housed in laminar flow racks and checked biweekly for viral infections (mouse hepatitis virus, minute virus of mice, rodent orphan parvovirus, Sendai) with kits obtained from Oreganon Teknika (Durham, NC) using established enzyme-linked immunoabsorbent assays. Mice testing positive for any of the above were omitted from the study. All mice were fed standard laboratory chow (Upjohn Lab Rodent Irradiated Mouse Chow, #5011-3, PMI Feeds, St. Louis, MO) and acidified drinking water (pH 5.0) ad libitum.

Induction of Inflammation by Intrapleural Injection of Dextran:

Intrapleural injections were made using a 27G needle cut to 3-4 mm and blunted by filing. Injections were made by inserting the needle between the mid-intercostal ribs on the right side of the thoracic cavity.

Dextran (MW 5-40x10⁶, St Louis, MO.) was injected as a 10% solution in saline in a volume of 100 µl/mouse. Care was taken to avoid bleeding at the site of injection at which the intercostal muscles were cut to facilitate smooth insertion of the needle.

Quantitation of Pleural Inflammatory Leukocyte Responses:

Pleural leukocytes were collected as follows: 4h post-induction, pleural inflammatory exudate was removed by washing with 2 x 1.0 ml Ca⁺⁺/Mg⁺⁺ free HBSS (Gibco, Grand Island, NY) containing 45 mg EDTA/100 ml HBSS, 40°C. Total leukocyte counts were made by hemocytometer following erythrocyte lysis in 2% acetic acid in PBS buffer; exudate leukocyte pellets were resuspended in serum for cytospin preparations and stained (Diff Quik, Baxter Healthcare, McGraw Park, IL) for differential leukocyte counts (neutrophils, eosinophils, and mononuclear leukocytes).

The pleural cavities of mice receiving either no intrapleural injection, or saline were washed and the cells counted in the same way to estimate baseline or saline-induced pleural leukocyte counts respectively.

Administration of compounds:

All drugs were dissolved in PBS and the pH adjusted to 7.5 with NaOH. Each compound was administered intravenously through the retroorbital sinus at hourly intervals (0-3h) starting from time "0" as indicated. Mice were carefully monitored for side effects; none were noted for the series of compounds reported herein.

The following compounds were tested for their inhibitory effects on dextran-induced leukocyte infiltration: Examples 3, 8, 9, 10, 12, 16, 37, 62, 66, 67, 99, 100, 111, 113, 115, 127, 131, 141, 184, 185, 192, PBS (saline) was administered iv. as a control.

Inhibition of eosinophil infiltration, which was suppressed by anti-alpha-4 Mab (PS/2, 50%), was used as a readout of VLA-4 antagonist activity of the compounds tested. Data for neutrophils are also reported.

Results:

- 5 Dextran pleural leukocyte response. The total pleural leukocyte counts were 255×10^4 (± 16 SEM) cells in the normal pleural cavity; of the normal pleural leukocyte population, all cells were mononuclear (a similar response was observed following intrapleural saline injection). Four hours after intrapleural injection of dextran total pleural leukocyte counts increased to 719×10^4 (± 67 SEM) and comprised 36.8×10^4 (± 4.1 SEM) eosinophils, 10 292×10^4 (± 25 SEM) neutrophils and 391×10^4 (± 48 SEM) mononuclear leukocytes.

% inhibition of Eosinophil infiltration

A > 40; B: 20-39; C < 19

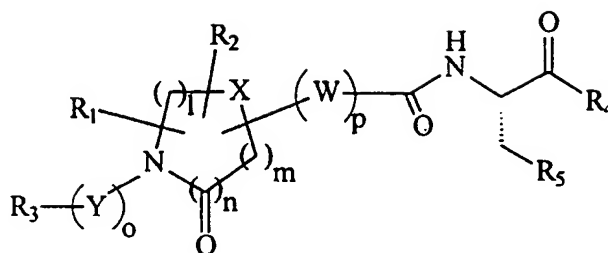
Example	Dose	Eos
3	50 x 2 iv	A
8	100 x 1 po	C
9	50 x 2 iv	A
10	100 x 1 po	B
12	50 x 2 iv	A
16	50 x 2 iv	A
37	50 x 2 iv	A
62	50 x 2 iv	B
66	50 x 2 iv	A
67	50 x 2 iv	A
99	50 x 2 iv	A
100	50 x 2 iv	B
111	50 x 2 iv	C
113	50 x 2 iv	C

Example	Dose	Eos
115	50 x 2 iv	B
127	50 x 2 iv	B
131	50 x 2 iv	C
141	50 x 2 iv	A
185	50 x 2 iv	B
184	50 x 2 iv	B
192	50 x 2 iv	B

CLAIMS

What is claimed is:

1. A compound of the formula:

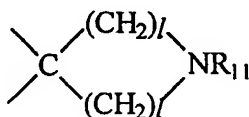


wherein

R_1 may occur one to four times and each occurrence is independently hydrogen or C_{1-6} alkyl;

- 5 R_2 is hydrogen, pyridyl, C_{1-6} alkyl, $(C_{1-6}$ alkyl)- CO_2-R_{11} , or $-CO_2-R_{11}$;

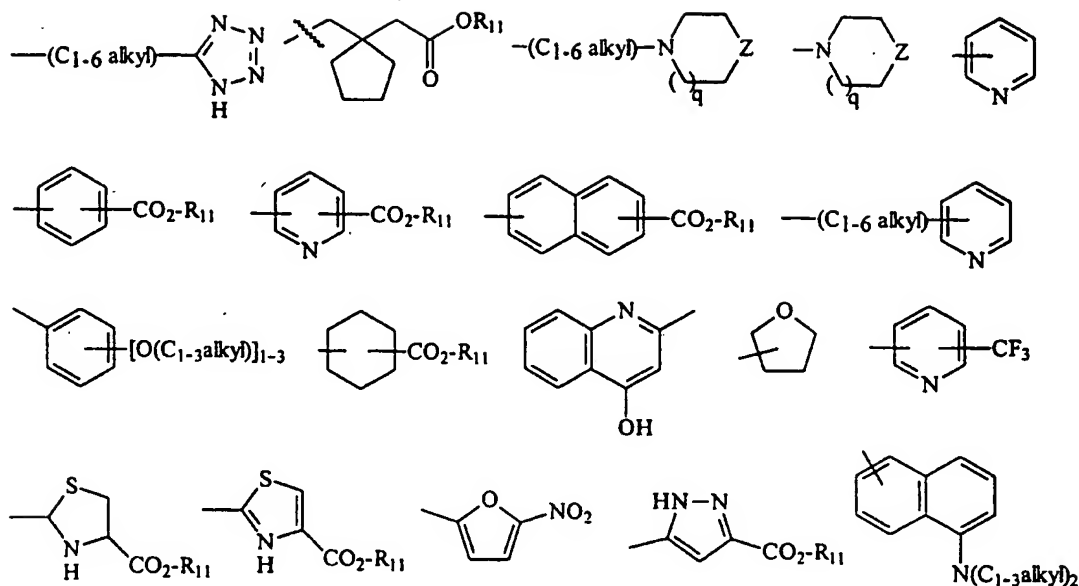
in addition, R_1 and R_2 may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



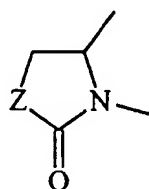
- 10 together with the carbon atom to which they are attached:

R_3 is hydrogen, phenyl, C_{1-6} alkyl, C_{3-6} alkenyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)- CO_2-R_{11} , $(C_{2-6}$ alkenyl)- CO_2-R_{11} , $(C_{1-6}$ alkyl)- $CO-C_{1-6}$ alkyl, $(C_{1-6}$ alkyl)- $O-C_{1-6}$ alkyl, $(C_{1-6}$ alkyl)-OH, $(C_{1-6}$ alkyl)-CN, adamantyl or one of the following:

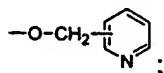
-266-



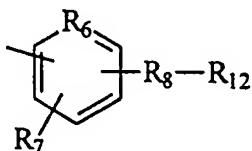
- 15 in addition, R_2 and R_3 -(Y)_o- may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:



R_4 is $-O-R_{11}$, NH_2 , $NHOH$, $-O-(C_{7-10} \text{ arylalkyl})$, or is of the formula



R_5 is a formula of the following:



20

R_6 is N or CH;

R_7 is hydrogen or halogen;

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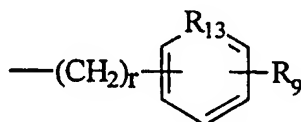
R_8 is $-NH-Y_1-$, $-OCH_2-$ or $-CONH-$;

R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or
 25 trifluoromethyl;

R_{10} is C_{1-6} alkyl, or $(C_{1-6} \text{ alkyl})-OH$, or hydrogen;

R_{11} is hydrogen or C_{1-6} alkyl;

R_{12} is C_{1-6} alkyl or the following formula:



30 R_{13} is N or CH;

W is $(C_{1-6} \text{ alkyl})$;

X is S, O, or CH_2 ;

Y and Y_1 are independently $-CO-$, $-C(=O)O-$, $-SO_2-$, or $-C(=O)N(R_{10})-$;

Z is O, CH_2 , or $N-R_{11}$;

35 l is 1, 2, or 3;

m is 1 or 2;

n is 0 or 1;

o is 0 or 1;

p is 0 or 1;

40 q is 0 or 1; and

r is 0, 1, 2 or 3;

with the provisos that

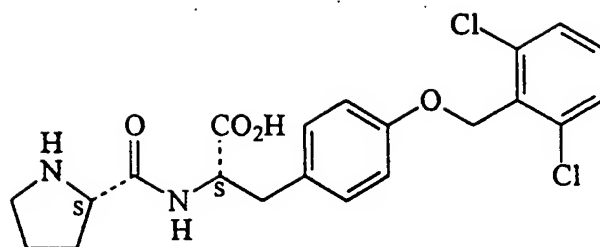
(1) when Y is $-C(=O)O-$, R_3 cannot be hydrogen;

(2) when R_4 is equal to $O-(C_4 \text{ alkyl})$, C_4 alkyl is not equal to *tert*-butyl;

45 (3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is CH_2), W is equal to CH_2 ; and

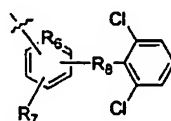
(4) the compound is not

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2. The compound according to claim 1,
wherein

R_3 is a formula of the following:



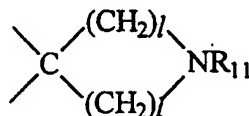
wherein R_7 is hydrogen or Cl.

3. The compound according to claim 2,
wherein

R_2 is hydrogen or C_{1-6} alkyl;

in addition, R_1 and R_2 may be attached to the same carbon atom and may form a

- 5 carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached or
be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



together with the carbon atom to which they are attached;

n is 0;

m is 2; and

p is 0.

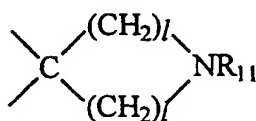
-269-

4. The compound according to claim 3,
wherein

R_1 is hydrogen or C_{1-3} alkyl;

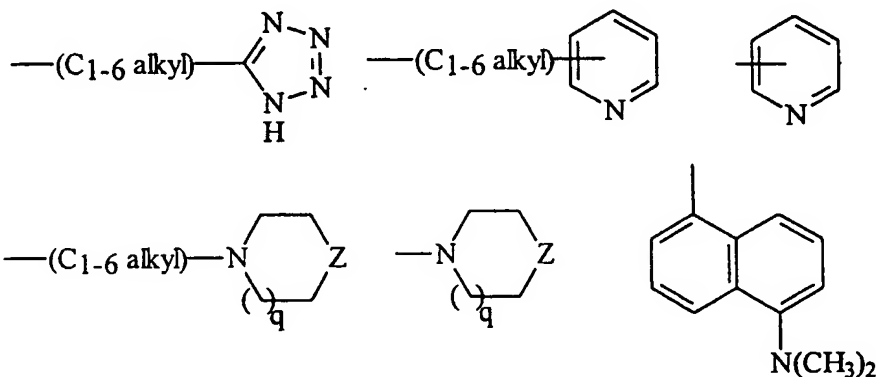
R_2 is hydrogen or C_{1-4} alkyl;

- 5 in addition, R_1 and R_2 may be attached to the same carbon atom and may form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached or be attached to the same atom and form a ring of 5-8 atoms of the formula:

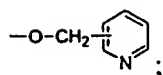


together with the carbon atom to which they are attached;

- 10 R_3 is hydrogen, C_{1-6} alkyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)-OH, $(C_{1-6}$ alkyl)-CO₂-R₁₁, $(C_{1-6}$ alkyl)-CN, adamantyl, phenyl, or one of the following:



R_4 is -O-R₁₁, NH₂, NHOH, or is of the formula



R_{11} is hydrogen or CH₃; and

X is S or O.

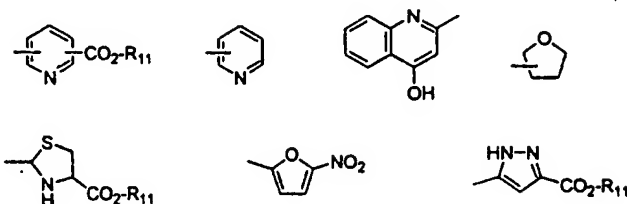
5. The compound according to claim 3,
wherein

R_1 is hydrogen;

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R_2 is hydrogen;

- 5 R_3 is $(C_{2-6}$ alkenyl)- CO_2-R_{11} , $(C_{1-6}$ alkyl)-O- C_{1-3} alkyl, $(C_{1-6}$ alkyl)- CO_2R_{11} , or one of the following:



R_4 is O- R_{11} ;

R_6 is CH;

R_{11} is hydrogen;

- 10 R_7 is hydrogen;

X is S;

Y is -CO-; and

l is 1.

6. The compound according to claim 3,

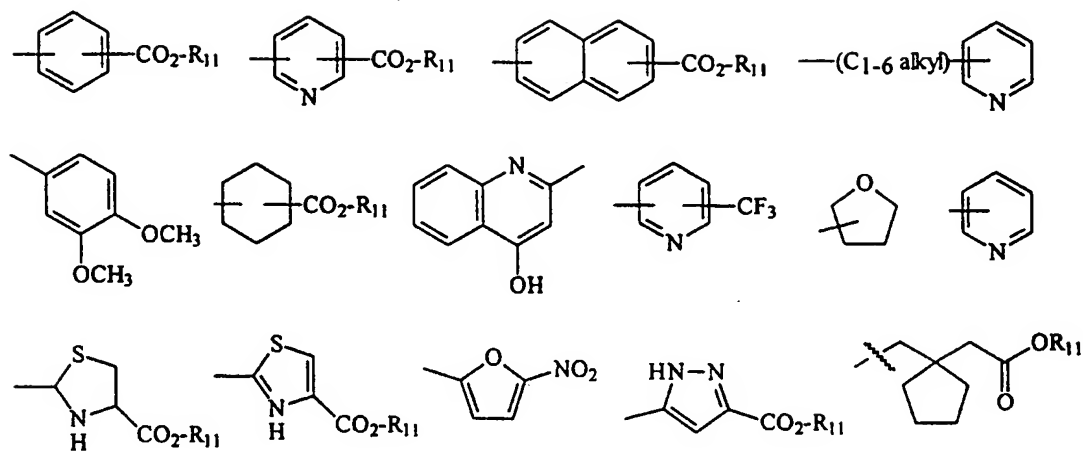
wherein

R_1 is hydrogen;

R_2 is hydrogen;

- 5 R_3 is C_{1-6} alkyl, $(C_{1-6}$ alkyl)- CO_2-R_{11} , $(C_{2-6}$ alkenyl)- CO_2-R_{11} , $(C_{1-6}$ alkyl)-CO- C_{1-6} alkyl, $(C_{1-6}$ alkyl)-O- C_{1-3} alkyl, $(C_{1-6}$ alkyl)-CN, or one of the following:

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R_4 is OH;

R_6 is CH;

R_{11} is hydrogen;

10

R_7 is hydrogen;

X is CH_2 ; and

Y is $-\text{CO}-$ or $-\text{C}(=\text{O})\text{NH}-$.

7. The compound according to claim 2,

wherein

W is $(\text{C}_{1-3} \text{ alkyl})$;

X is CH_2 ;

5

Y is $-\text{C}(=\text{O})\text{O}-$;

R_1 is hydrogen;

R_2 is hydrogen, $(\text{C}_{1-3} \text{ alkyl})-\text{CO}_2-\text{R}_{11}$, or $-\text{CO}_2-\text{R}_{11}$;

R_3 is hydrogen, C_{7-10} arylalkyl, C_{1-6} alkyl, or $(\text{C}_{1-6} \text{ alkyl})-\text{CO}_2-\text{R}_{11}$;

R_4 is OH;

10

R_6 is CH;

R_{11} is hydrogen;

R_7 is hydrogen;

l is 1 or 3; and

-272-

n is 0.

8. The compound according to claim 2,
wherein

W is C₁₋₃ alkyl;

X is CH₂;

5 R₁ is hydrogen;

R₂ is (C₁₋₄ alkyl)-CO₂-R₁₁, or CO₂-R₁₁;

R₃ is hydrogen, C₁₋₃ alkyl, or C₂₋₇ alkenyl;

R₄ is OH,

R₆ is CH;

10 R₁₁ is hydrogen;

R₇ is hydrogen;

l is 1;

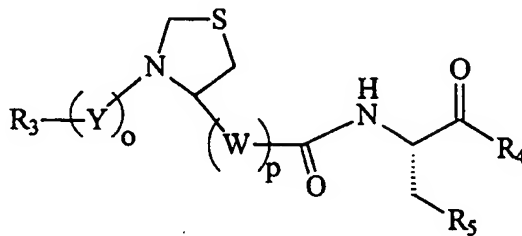
m is 1;

n is 1;

15 o is 0; and

p is 1.

9. The compound according to claim 1, wherein said compound is



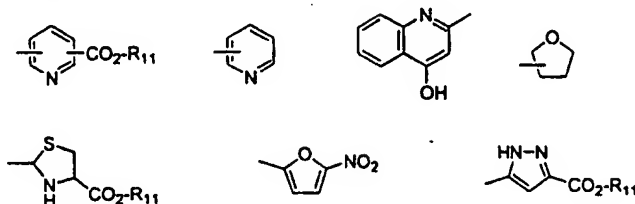
10. The compound according to claim 4,
wherein

R₁ is hydrogen;

-273-

R_2 is hydrogen;

- 5 R_3 is C_{1-6} alkyl, $(C_{1-6} \text{ alkyl})-CO_2R_{11}$, $(C_{2-6} \text{ alkenyl})-CO_2R_{11}$, $(C_{1-6} \text{ alkyl})-O-C_{1-3} \text{ alkyl}$,
or one of the following:



R_4 is $O-R_{11}$;

R_6 is CH ;

R_{11} is hydrogen or C_{1-6} alkyl;

- 10 R_7 is hydrogen;

X is S ;

Y is $-C(=O)O-$; and

l is 1.

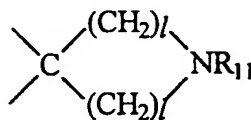
11. The compound according to claim 1,

wherein

R_1 is hydrogen or C_{1-3} alkyl;

R_2 is hydrogen or C_{1-4} alkyl;

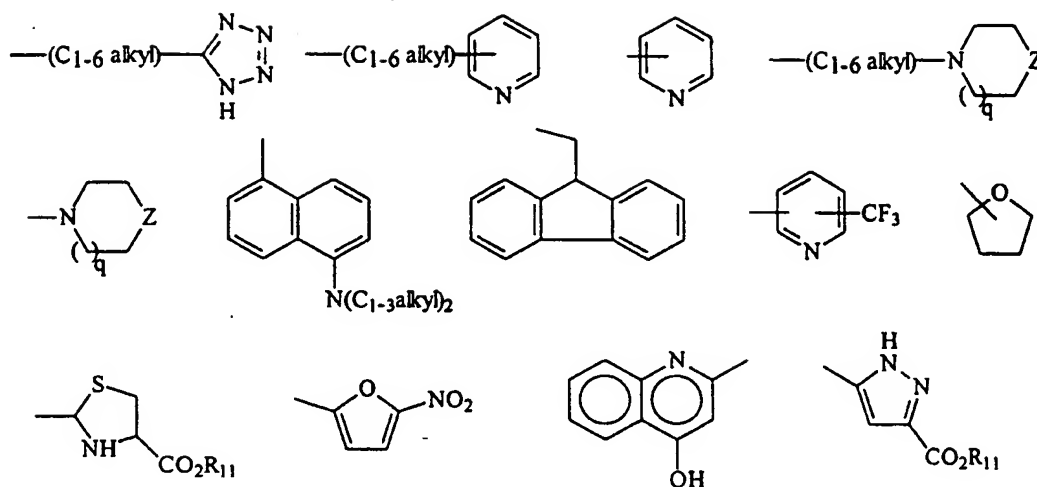
- 5 in addition, R_1 and R_2 may be attached to the same carbon atom and may form a
carbocyclic ring of 5-8 atoms, or be attached to the same atom and form a ring of 5-8
atoms of the formula:



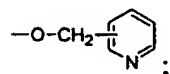
together with the carbon atom to which they are attached;

- 10 R_3 is hydrogen, C_{1-6} alkyl, C_{7-17} arylalkyl, $(C_{1-6} \text{ alkyl})-OH$, $(C_{1-6} \text{ alkyl})-CO_2R_{11}$, $(C_{1-6} \text{ alkyl})-CN$, adamantyl, phenyl, or one of the following:

-274-



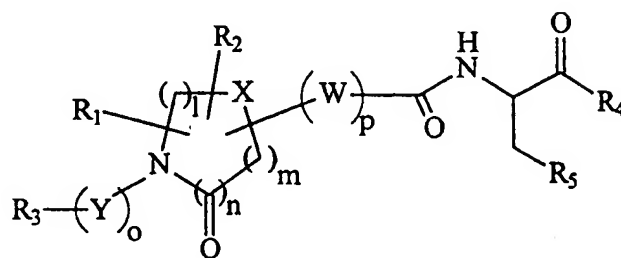
R_4 is $-O-R_{11}$, NH_2 , $NHOH$, or is of the formula



R_{11} is hydrogen or C_{1-6} alkyl; and

X is S or O.

12. A compound of the formula:

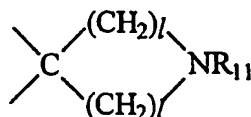


wherein

R_1 may occur one to four times and each occurrence is independently hydrogen or C_{1-6} alkyl;

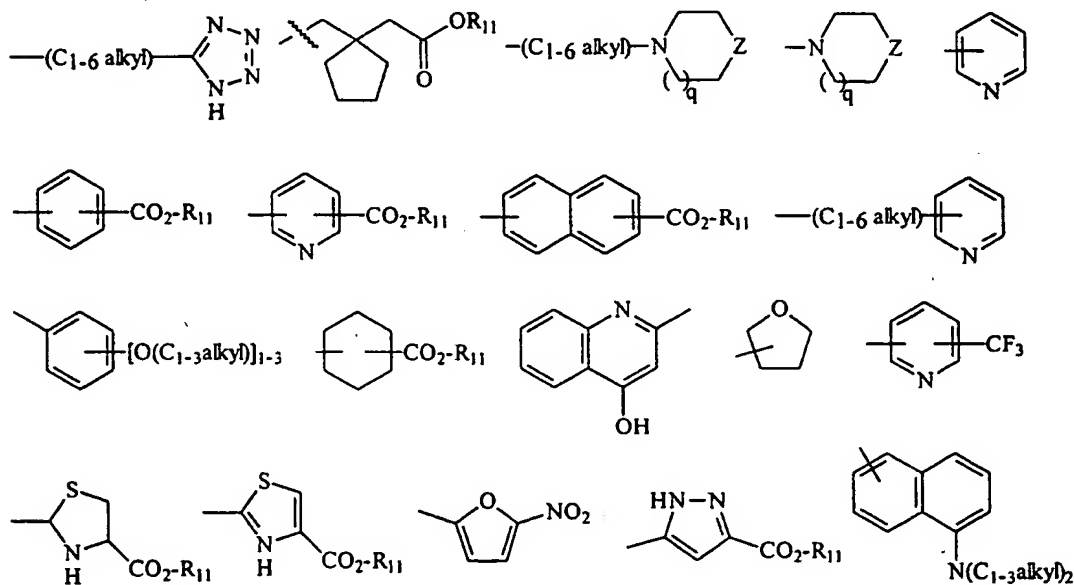
5 R_2 is hydrogen, pyridyl, C_{1-6} alkyl, $(C_{1-6} \text{ alkyl})-CO_2-R_{11}$, or $-CO_2-R_{11}$;

in addition, R_1 and R_2 may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:

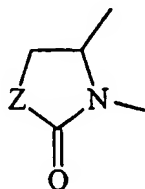


together with the carbon atom to which they are attached;

- 10 R_3 is hydrogen, phenyl, C_{1-6} alkyl, C_{3-6} alkenyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)- CO_2 - R_{11} , $(C_{2-6}$ alkenyl)- CO_2 - R_{11} , $(C_{1-6}$ alkyl)- CO - C_{1-6} alkyl, $(C_{1-6}$ alkyl)- O - C_{1-6} alkyl, $(C_{1-6}$ alkyl)- OH , $(C_{1-6}$ alkyl)- CN , adamantyl or one of the following:

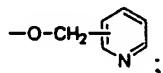


in addition, R_2 and $R_3-(Y)_o$ may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:

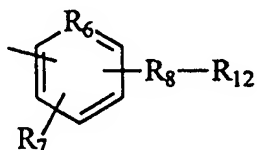


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R_4 is $-O-R_{11}$, NH_2 , $NHOH$, $-O-(C_{7-10} \text{ arylalkyl})$, or is of the formula



R_5 is a formula of the following:



R_6 is N or CH;

20 R_7 is hydrogen or halogen;

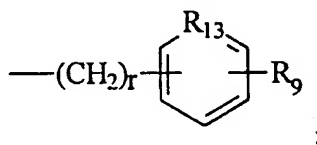
R_8 is $-NH-Y_1-$, $-OCH_2-$ or $-CONH-$;

R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or trifluoromethyl;

R_{10} is C_{1-6} alkyl, or $(C_{1-6} \text{ alkyl})-OH$, or hydrogen;

25 R_{11} is hydrogen or C_{1-6} alkyl;

R_{12} is C_{1-6} alkyl or the following formula:



R_{13} is N or CH;

W is $(C_{1-6} \text{ alkyl})$;

30 X is S, O, or CH_2 ;

Y and Y_1 are independently $-CO-$, $-C(=O)O-$, $-SO_2-$, or $-C(=O)N(R_{10})-$;

Z is O, CH_2 , or $N-R_{11}$;

l is 1, 2, or 3;

m is 1 or 2;

35 n is 0 or 1;

o is 0 or 1;

p is 0 or 1;

q is 0 or 1; and

-277-

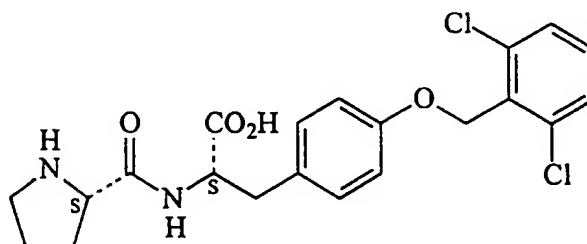
r is 0, 1, 2 or 3;

40 with the provisos that

- (1) when Y is $-C(=O)O-$, R_3 cannot be hydrogen;
- (2) when R_4 is equal to $O-(C_4 \text{ alkyl})$, C_4 alkyl is not equal to *tert*-butyl;
- (3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is CH_2),

W is equal to CH_2 ;

- 45 (4) the compound has an IC_{50} value of less than $5\mu M$ in a Jurkat CS-1 assay and/or an IC_{50} value of less than $50\mu M$ in a Jurkat EC assay; and
- (5) the compound is not



13. A pharmaceutical composition comprising:

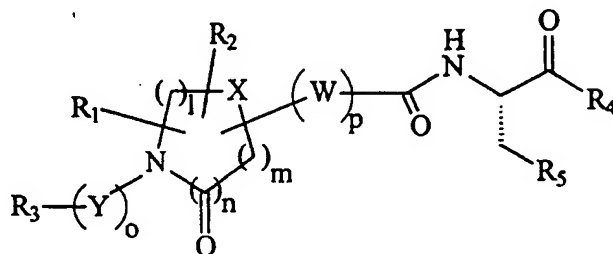
- a therapeutically effective amount of the compound as set forth in any one of the preceding claims; and
- a pharmaceutically acceptable carrier or diluent.

14. A method for treating or preventing $\alpha_4\beta_1$ adhesion mediated conditions in a human which comprises administering to a patient an effective amount of the compound according to any one of claims 1, 2, 3, 4, 5, 6, 7, 8, 9, 10, 11, or 12.

15. The method according to claim 14, wherein said condition is selected from the group consisting of rheumatoid arthritis, asthma, allergy conditions, allograft rejection, psoriasis, eczema, contact dermatitis and other skin inflammatory diseases, inflammatory and immunoinflammatory conditions including ophthalmic inflammatory conditions,
5 inflammatory bowel diseases, atherosclerosis, and ulcerative colitis.

16. The method according to claim 15, wherein said condition is asthma.

17. A compound of the formula:

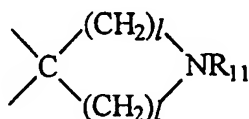


wherein

R_1 may occur one to four times and each occurrence is independently hydrogen or C_{1-6} alkyl;

5 R_2 is hydrogen, pyridyl, C_{1-6} alkyl, $(C_{1-6} \text{ alkyl})-CO_2-R_{11}$, or $-CO_2-R_{11}$;

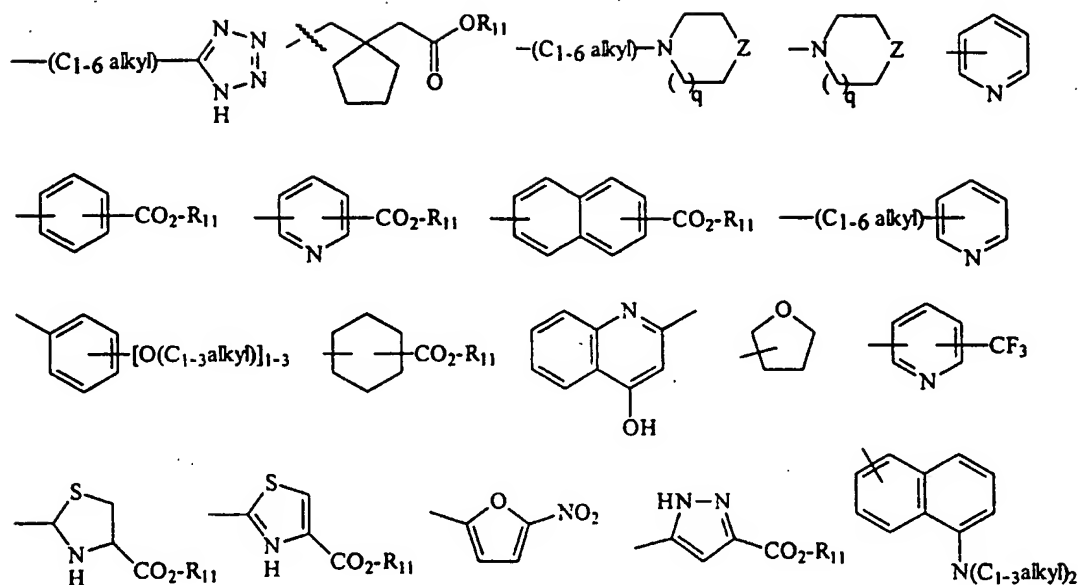
in addition, R_1 and R_2 may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



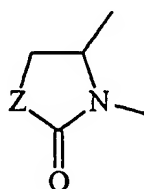
together with the carbon atom to which they are attached;

10 R_3 is hydrogen, phenyl, C_{1-6} alkyl, C_{3-6} alkenyl, C_{7-17} arylalkyl, $(C_{1-6} \text{ alkyl})-CO_2-R_{11}$, $(C_{2-6} \text{ alkenyl})-CO_2-R_{11}$, $(C_{1-6} \text{ alkyl})-CO-C_{1-6} \text{ alkyl}$, $(C_{1-6} \text{ alkyl})-O-C_{1-6} \text{ alkyl}$, $(C_{1-6} \text{ alkyl})-OH$, $(C_{1-6} \text{ alkyl})-CN$, adamantyl or one of the following:

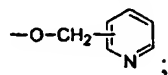
-279-



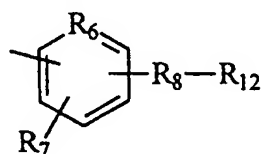
in addition, R_2 and R_3 -(Y)_o- may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:



R_4 is $-O-R_{11}$, NH_2 , $NHOH$, $-O-(C_{7-10} \text{ arylalkyl})$, or is of the formula



R_5 is a formula of the following:



R_6 is N or CH;

R_7 is hydrogen or halogen;

R_8 is -NHCO- ;

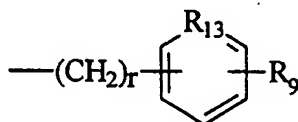
R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or trifluoromethyl;

R_{10} is C_{1-6} alkyl, or $(\text{C}_{1-6} \text{ alkyl})\text{-OH}$, or hydrogen;

25

R_{11} is hydrogen or C_{1-6} alkyl;

R_{12} is C_{1-6} alkyl or the following formula:



R_{13} is N or CH;

W is $(\text{C}_{1-6} \text{ alkyl})$;

30

X is S, O, or CH_2 ;

Y is -CO- , -C(=O)O- , or $\text{-C(=O)N(R}_{10}\text{)-}$;

Z is O, CH_2 , or N-R_{11} ;

l is 1, 2, or 3;

m is 1 or 2;

35

n is 0 or 1;

o is 0 or 1;

p is 0 or 1;

q is 0 or 1; and

r is 0, 1, 2 or 3;

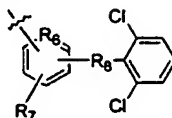
40

with the provisos that

- (1) when Y is -C(=O)O- , R_3 cannot be hydrogen;
- (2) when R_4 is equal to $\text{O-(C}_4 \text{ alkyl)}$, $\text{C}_4 \text{ alkyl}$ is not equal to *tert*-butyl; and
- (3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is CH_2), W is equal to CH_2 .

18. The compound according to claim 17,
wherein

R_3 is a formula of the following:

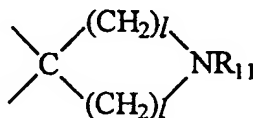


wherein R_7 is hydrogen or Cl.

19. The compound according to claim 18,
wherein

R_2 is hydrogen or C_{1-6} alkyl;

in addition, R_1 and R_2 may be attached to the same carbon atom and may form a
5 carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached or
be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



together with the carbon atom to which they are attached;

n is 0;

m is 2; and

10 p is 0.

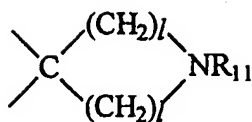
20. The compound according to claim 19,
wherein

R_1 is hydrogen or C_{1-3} alkyl;

R_2 is hydrogen or C_{1-4} alkyl;

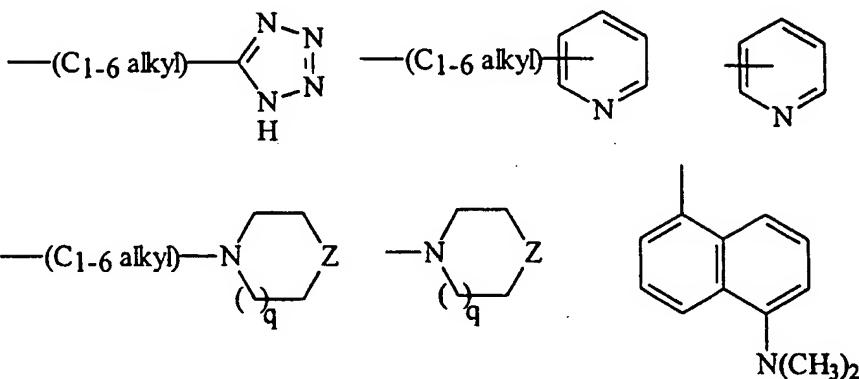
5 in addition, R_1 and R_2 may be attached to the same carbon atom and may form a
carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached or
be attached to the same atom and form a ring of 5-8 atoms of the formula:

-282-

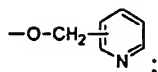


together with the carbon atom to which they are attached;

R_3 is hydrogen, C_{1-6} alkyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)-OH, $(C_{1-6}$ alkyl)-CO₂-R₁₁, $(C_{1-6}$ alkyl)-CN, adamantyl, phenyl, or one of the following:



R_4 is -O-R₁₁, NH₂, NHOH, or is of the formula



R_{11} is hydrogen or CH₃; and

X is S or O.

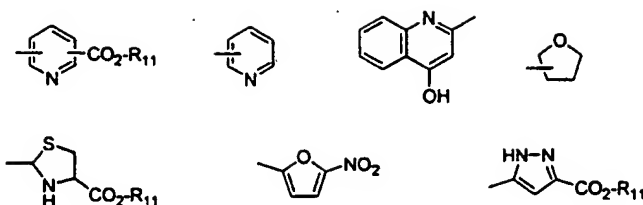
21. The compound according to claim 19,
wherein

R_1 is hydrogen;

R_2 is hydrogen;

5 R_3 is $(C_{2-6}$ alkenyl)-CO₂-R₁₁, $(C_{1-6}$ alkyl)-O-C₁₋₃ alkyl, $(C_{1-6}$ alkyl)-CO₂-R₁₁, or one of the following:

-283-



R_4 is $O-R_{11}$;

R_6 is CH ;

R_{11} is hydrogen;

10

R_7 is hydrogen;

X is S ;

Y is $-CO-$; and

l is 1.

22. The compound according to claim 19,

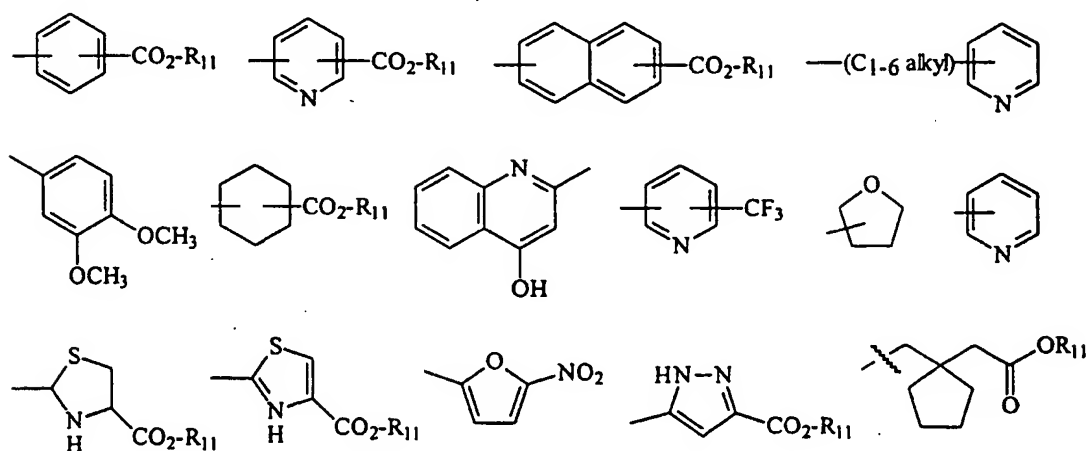
wherein

R_1 is hydrogen;

R_2 is hydrogen;

5

R_3 is C_{1-6} alkyl, $(C_{1-6}$ alkyl)- CO_2-R_{11} , $(C_{2-6}$ alkenyl)- CO_2-R_{11} , $(C_{1-6}$ alkyl)- $CO-C_{1-6}$ alkyl, $(C_{1-6}$ alkyl)- $O-C_{1-3}$ alkyl, $(C_{1-6}$ alkyl)- CN , or one of the following:



R_4 is OH ;

R_6 is CH ;

10 R₁₁ is hydrogen;
 R₇ is hydrogen;
 X is CH₂; and
 Y is -CO- or -C(=O)NH-.

23. The compound according to claim 18,
 wherein
 W is (C₁₋₃ alkyl);
 X is CH₂;
5 Y is C(=O)O-;
 R₁ is hydrogen;
 R₂ is hydrogen, (C₁₋₃ alkyl)-CO₂-R₁₁, or -CO₂-R₁₁;
 R₃ is hydrogen, C₇₋₁₀ arylalkyl, C₁₋₆ alkyl, or (C₁₋₆ alkyl)-CO₂-R₁₁;
 R₄ is OH;
10 R₆ is CH;
 R₁₁ is hydrogen;
 R₇ is hydrogen;
 l is 1 or 3; and
 n is 0.

24. The compound according to claim 18,
 wherein
 W is C₁₋₃ alkyl;
 X is CH₂;
5 R₁ is hydrogen;
 R₂ is (C₁₋₄ alkyl)-CO₂-R₁₁, or CO₂-R₁₁;
 R₃ is hydrogen, C₁₋₃ alkyl, or C₂₋₇ alkenyl;
 R₄ is OH,
 R₆ is CH;
10 R₁₁ is hydrogen;

-285-

R_7 is hydrogen;

l is 1;

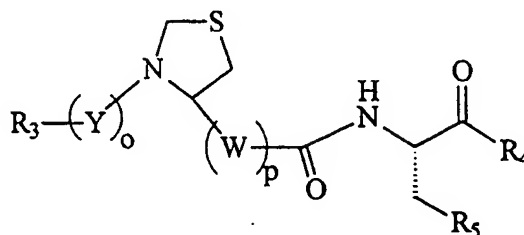
m is 1;

n is 1;

15 o is 0; and

p is 1.

25. The compound according to claim 17, wherein said compound is



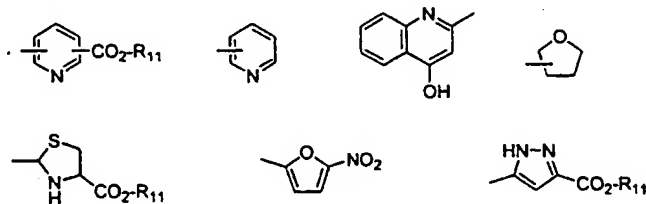
26. The compound according to claim 20,

wherein

R_1 is hydrogen;

R_2 is hydrogen;

5 R_3 is C_{1-6} alkyl, $(C_{1-6}$ alkyl)- CO_2R_{11} , $(C_{2-6}$ alkenyl)- CO_2R_{11} , $(C_{1-6}$ alkyl)-O- C_{1-3} alkyl, or one of the following:



R_4 is O- R_{11} ;

R_6 is CH;

R_{11} is hydrogen or C_{1-6} alkyl;

10 R_7 is hydrogen;

X is S;

Y is -C(=O)O-; and

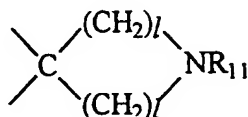
l is 1.

27. The compound according to claim 17,
wherein

R₁ is hydrogen or C₁₋₃ alkyl;

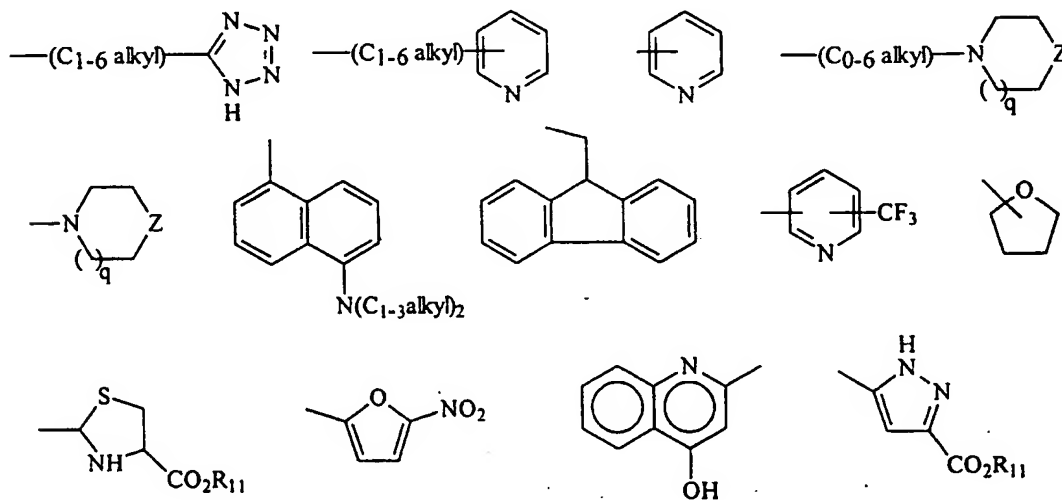
R₂ is hydrogen or C₁₋₄ alkyl;

5 in addition, R₁ and R₂ may be attached to the same carbon atom and may form a carbocyclic ring of 5-8 atoms, or be attached to the same atom and form a ring of 5-8 atoms of the formula:

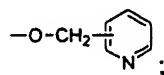


together with the carbon atom to which they are attached;

R₃ is hydrogen, C₁₋₆ alkyl, C₇₋₁₇ arylalkyl, (C₁₋₆ alkyl)-OH, (C₁₋₆ alkyl)-CO₂-R₁₁, (C₁₋₆ alkyl)-CN, adamantyl, phenyl, or one of the following:



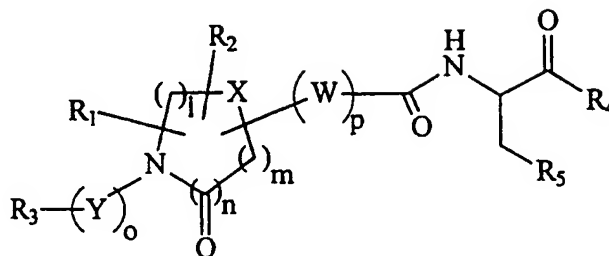
R₄ is -O-R₁₁, NH₂, NHOH, or is of the formula



R_{11} is hydrogen or C_{1-6} alkyl; and

X is S or O.

28. A compound of the formula:

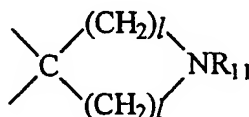


wherein

R_1 may occur one to four times and each occurrence is independently hydrogen or C_{1-6} alkyl;

5 R_2 is hydrogen, pyridyl, C_{1-6} alkyl, $(C_{1-6}$ alkyl)- CO_2-R_{11} , or $-CO_2-R_{11}$;

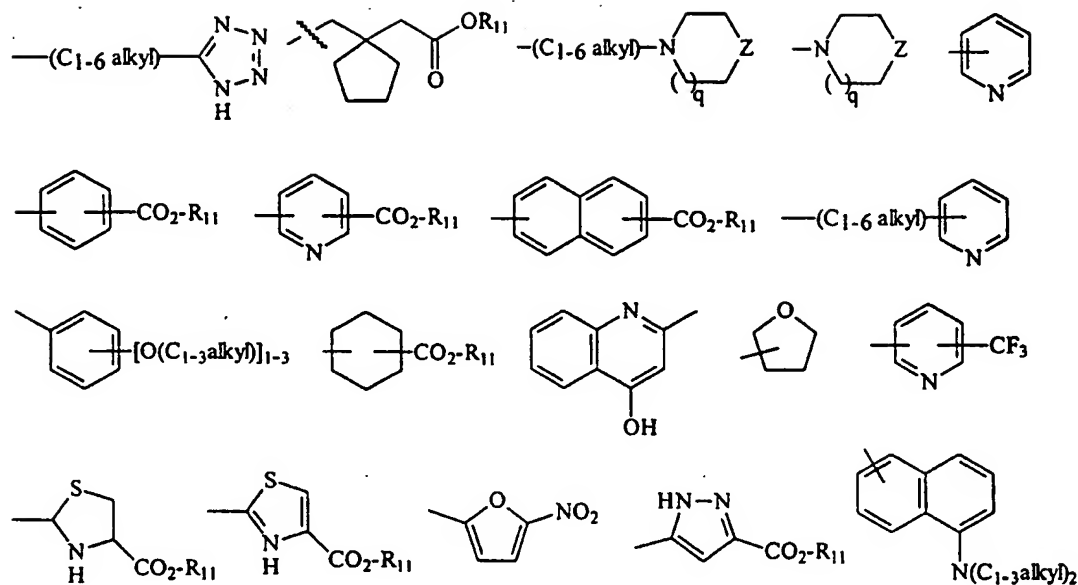
in addition, R_1 and R_2 may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



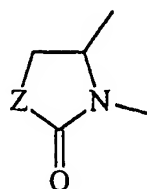
10 together with the carbon atom to which they are attached;

R_3 is hydrogen, phenyl, C_{1-6} alkyl, C_{3-6} alkenyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)- CO_2-R_{11} , $(C_{2-6}$ alkenyl)- CO_2-R_{11} , $(C_{1-6}$ alkyl)- $CO-C_{1-6}$ alkyl, $(C_{1-6}$ alkyl)- $O-C_{1-6}$ alkyl, $(C_{1-6}$ alkyl)-OH, $(C_{1-6}$ alkyl)-CN, adamantyl or one of the following:

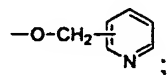
-288-



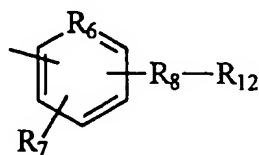
- 15 in addition, R_2 and R_3 -(Y)₆- may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:



R_4 is $-O-R_{11}$, NH_2 , $NHOH$, $-O-(C_{7-10} \text{ arylalkyl})$, or is of the formula



R_5 is a formula of the following:



20

R_6 is N or CH;

R_7 is hydrogen or halogen;

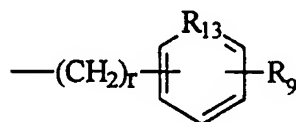
R_8 is -NHCO- ;

R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or trifluoromethyl;

25 R_{10} is C_{1-6} alkyl, or $(\text{C}_{1-6} \text{ alkyl})\text{-OH}$, or hydrogen;

R_{11} is hydrogen or C_{1-6} alkyl;

R_{12} is C_{1-6} alkyl or the following formula:



R_{13} is N or CH;

30 W is $(\text{C}_{1-6} \text{ alkyl})$;

X is S, O, or CH_2 ;

Y is -CO- , -C(=O)O- , or $\text{-C(=O)N(R}_{10}\text{)-}$;

Z is O, CH_2 , or N-R_{11} ;

l is 1, 2, or 3;

35 m is 1 or 2;

n is 0 or 1;

o is 0 or 1;

p is 0 or 1;

q is 0 or 1; and

40 r is 0, 1, 2 or 3;

with the provisos that

(1) when Y is -C(=O)O- , R_9 cannot be hydrogen;

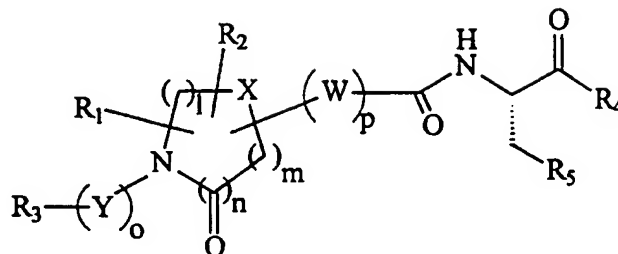
(2) when R_4 is equal to $\text{O-(C}_4 \text{ alkyl)}$, C_4 alkyl is not equal to *tert*-butyl;

(3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is

45 CH_2), W is equal to CH_2 ; and

(4) the compound has an IC_{50} value of less than $5\mu\text{M}$ in a Jurkat CS-1 assay and/or an IC_{50} value of less than $50\mu\text{M}$ in a Jurkat EC assay.

29. A compound of the formula:

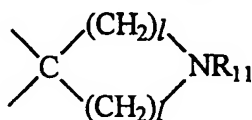


wherein

R₁ may occur one to four times and each occurrence is independently hydrogen or C₁₋₆ alkyl;

5 R₂ is hydrogen, pyridyl, C₁₋₆ alkyl, (C₁₋₆ alkyl)-CO₂-R₁₁, or -CO₂-R₁₁;

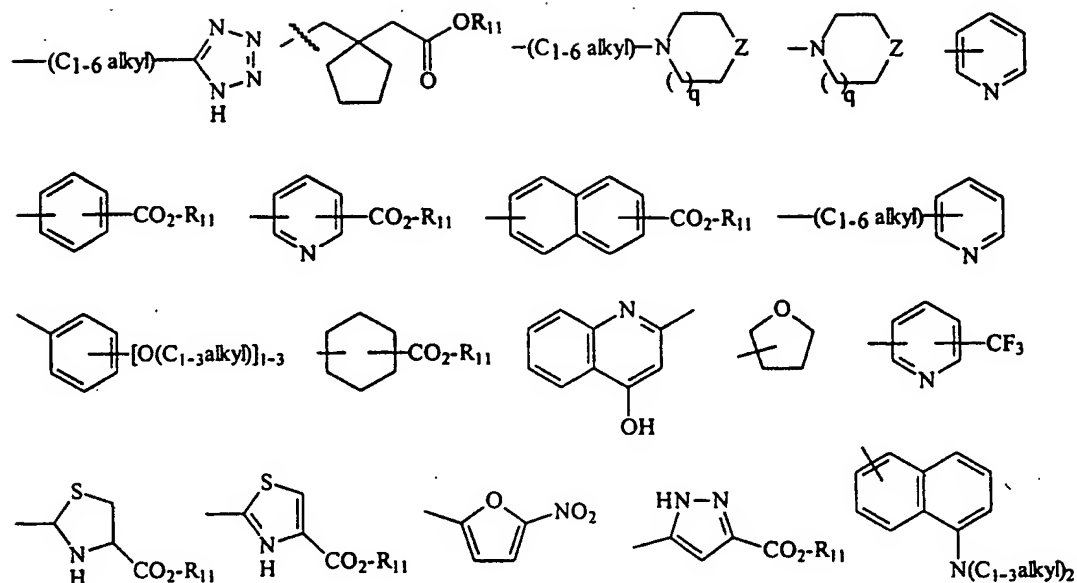
in addition, R₁ and R₂ may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



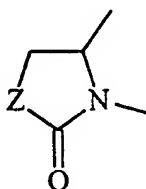
together with the carbon atom to which they are attached;

10 R₃ is hydrogen, phenyl, C₁₋₆ alkyl, C₃₋₆ alkenyl, C₇₋₁₇ arylalkyl, (C₁₋₆ alkyl)-CO₂-R₁₁, (C₂₋₆ alkenyl)-CO₂-R₁₁, (C₁₋₆ alkyl)-CO-C₁₋₆ alkyl, (C₁₋₆ alkyl)-O-C₁₋₆ alkyl, (C₁₋₆ alkyl)-OH, (C₁₋₆ alkyl)-CN, adamantyl or one of the following:

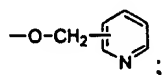
-291-



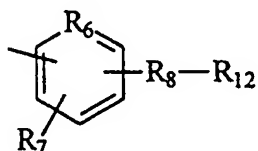
in addition, R_2 and R_3 -(Y)₆- may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom to which they are attached:



R_4 is $-O-R_{11}$, NH_2 , $NHOH$, $-O-(C_{7-10} \text{ arylalkyl})$, or is of the formula



R_5 is a formula of the following:



R_6 is N or CH;

R_7 is hydrogen or halogen;

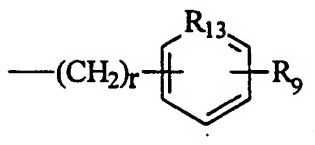
R_8 is $-\text{OCH}_2-$;

R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or trifluoromethyl;

R_{10} is C_{1-6} alkyl, or $(\text{C}_{1-6} \text{ alkyl})-\text{OH}$, or hydrogen;

25 R_{11} is hydrogen or C_{1-6} alkyl;

R_{12} is C_{1-6} alkyl or the following formula:



R_{13} is N or CH;

W is $(\text{C}_{1-6} \text{ alkyl})$;

30 X is S, O, or CH_2 ;

Y is $-\text{CO}-$, $-\text{C}(=\text{O})\text{O}-$, or $-\text{C}(=\text{O})\text{N}(\text{R}_{10})-$;

Z is O, CH_2 , or $\text{N}-\text{R}_{11}$;

l is 1, 2, or 3;

m is 1 or 2;

35 n is 0 or 1;

o is 0 or 1;

p is 0 or 1;

q is 0 or 1; and

r is 0, 1, 2 or 3;

40 with the provisos that

(1) when Y is $-\text{C}(=\text{O})\text{O}-$, R_3 cannot be hydrogen;

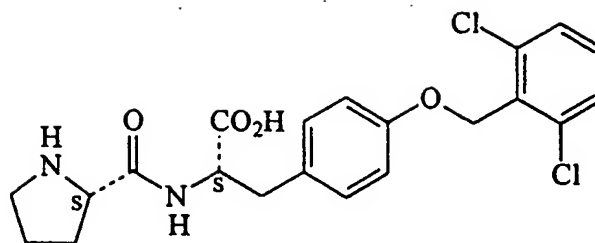
(2) when R_4 is equal to $\text{O}-(\text{C}_4 \text{ alkyl})$, $\text{C}_4 \text{ alkyl}$ is not equal to *tert*-butyl;

(3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is CH_2), W is equal to CH_2 ;

45 (4) when R_3 is phenyl, C_{1-6} alkyl, C_{7-17} arylalkyl, $(\text{C}_{1-6} \text{ alkyl})-\text{CO}_2-\text{R}_{11}$, $(\text{C}_{1-6} \text{ alkyl})-\text{O}-\text{C}_{1-6} \text{ alkyl}$, or $(\text{C}_{1-6} \text{ alkyl})\text{OH}$, o is 0; and

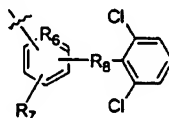
(5) the compound is not

-293-



30. The compound according to claim 29,
wherein

R_3 is a formula of the following:

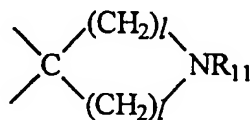


wherein R_7 is hydrogen or Cl.

31. The compound according to claim 30,
wherein

R_2 is hydrogen or C_{1-6} alkyl;

- in addition, R_1 and R_2 may be attached to the same carbon atom and may form a
5 carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached or
be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:



together with the carbon atom to which they are attached;

n is 0;

m is 2; and

p is 0.

-294-

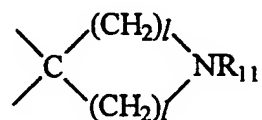
32. The compound according to claim 31,

wherein

R_1 is hydrogen or C_{1-3} alkyl;

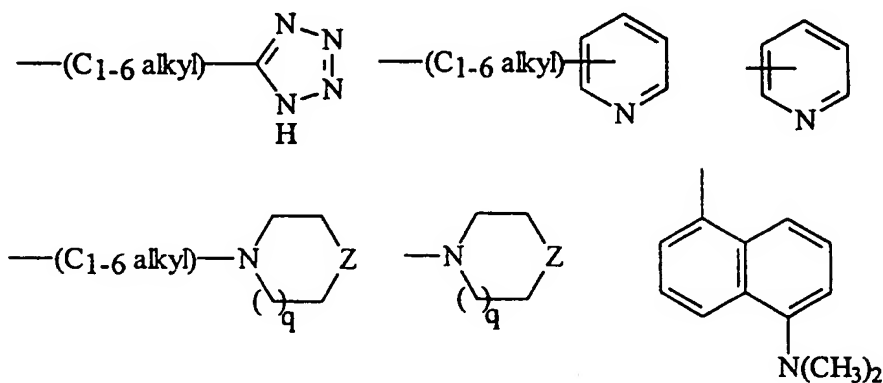
R_2 is hydrogen or C_{1-4} alkyl;

- 5 in addition, R_1 and R_2 may be attached to the same carbon atom and may form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached or be attached to the same atom and form a ring of 5-8 atoms of the formula:

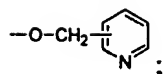


together with the carbon atom to which they are attached;

- 10 R_3 is hydrogen, C_{1-6} alkyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)-OH, $(C_{1-6}$ alkyl)-CO₂-R₁₁, $(C_{1-6}$ alkyl)-CN, adamantyl, phenyl, or one of the following:



R_4 is -O-R₁₁, NH₂, NHOH, or is of the formula



R_{11} is hydrogen or CH₃; and

X is S or O.

33. The compound according to claim 31,

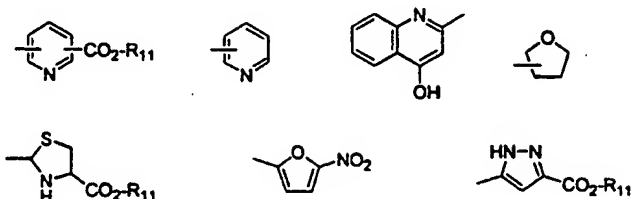
wherein

R_1 is hydrogen;

-295-

R_2 is hydrogen;

- 5 R_3 is $(C_{2-6}$ alkenyl)- CO_2-R_{11} , $(C_{1-6}$ alkyl)-O- C_{1-3} alkyl, $(C_{1-6}$ alkyl)- CO_2R_{11} , or one of the following:



R_4 is O- R_{11} ;

R_6 is CH;

R_{11} is hydrogen;

- 10 R_7 is hydrogen;

X is S;

Y is -CO-; and

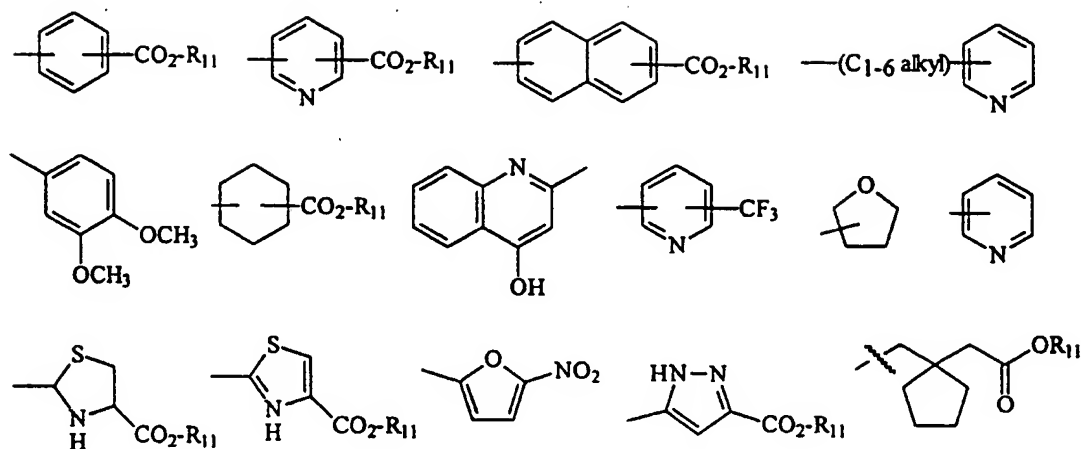
l is 1.

34. The compound according to claim 31,
wherein

R_1 is hydrogen;

R_2 is hydrogen;

- 5 R_3 is C_{1-6} alkyl, $(C_{1-6}$ alkyl)- CO_2-R_{11} , $(C_{2-6}$ alkenyl)- CO_2-R_{11} , $(C_{1-6}$ alkyl)-CO- C_{1-6} alkyl, $(C_{1-6}$ alkyl)-O- C_{1-3} alkyl, $(C_{1-6}$ alkyl)-CN, or one of the following:



R_4 is OH;

R_6 is CH;

R_{11} is hydrogen;

10

R_7 is hydrogen;

X is CH_2 ; and

Y is $-CO-$ or $-C(=O)NH-$.

35. The compound according to claim 30,

wherein

W is $(C_{1-3} \text{ alkyl})$;

X is CH_2 ;

5

Y is $-C(=O)O-$;

R_1 is hydrogen;

R_2 is hydrogen, $(C_{1-3} \text{ alkyl})-CO_2-R_{11}$, or $-CO_2-R_{11}$;

R_3 is hydrogen, C_{7-10} arylalkyl, C_{1-6} alkyl, or $(C_{1-6} \text{ alkyl})-CO_2-R_{11}$;

R_4 is OH;

10

R_6 is CH;

R_{11} is hydrogen;

R_7 is hydrogen;

l is 1 or 3; and

n is 0.

36. The compound according to claim 30,
wherein

W is C₁₋₃ alkyl;

X is CH₂;

5 R₁ is hydrogen;

R₂ is (C₁₋₄ alkyl)-CO₂-R₁₁, or CO₂-R₁₁;

R₃ is hydrogen, C₁₋₃ alkyl, or C₂₋₇ alkenyl;

R₄ is OH,

R₆ is CH;

10 R₁₁ is hydrogen;

R₇ is hydrogen;

l is 1;

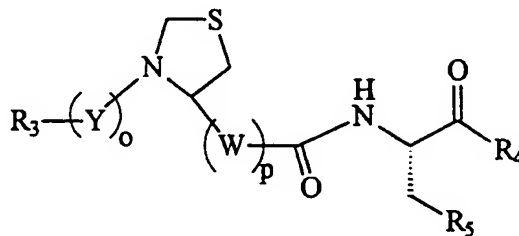
m is 1;

n is 1;

15 o is 0; and

p is 1.

37. The compound according to claim 29, wherein said compound is



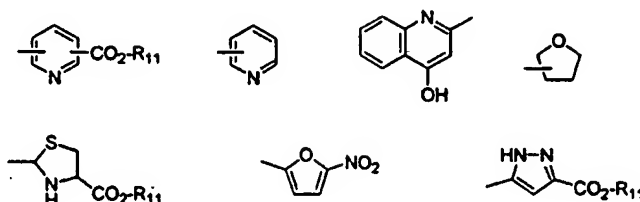
38. The compound according to claim 32,
wherein

R₁ is hydrogen;

-298-

R_2 is hydrogen;

- 5 R_3 is C_{1-6} alkyl, $(C_{1-6}$ alkyl)- CO_2R_{11} , $(C_{2-6}$ alkenyl)- CO_2R_{11} , $(C_{1-6}$ alkyl)-O- C_{1-3} alkyl, or one of the following:



R_4 is O- R_{11} ;

R_6 is CH;

R_{11} is hydrogen or C_{1-6} alkyl;

- 10 R_7 is hydrogen;

X is S;

Y is $-C(=O)O-$; and

l is 1.

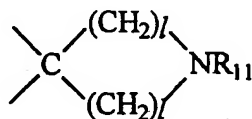
39. The compound according to claim 29,

wherein

R_1 is hydrogen or C_{1-3} alkyl;

R_2 is hydrogen or C_{1-4} alkyl;

- 5 in addition, R_1 and R_2 may be attached to the same carbon atom and may form a carbocyclic ring of 5-8 atoms, or be attached to the same atom and form a ring of 5-8 atoms of the formula:

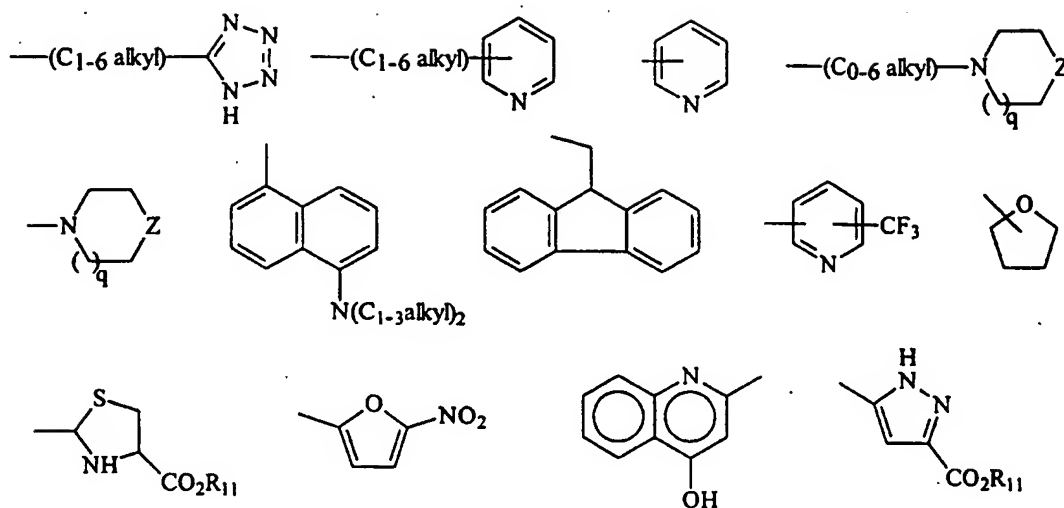


together with the carbon atom to which they are attached;

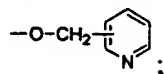
R_3 is hydrogen, C_{1-6} alkyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)-OH, $(C_{1-6}$ alkyl)- CO_2R_{11} ,

- 10 $(C_{1-6}$ alkyl)-CN, adamantyl, phenyl, or one of the following:

-299-



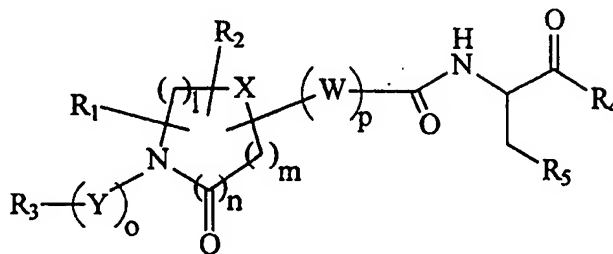
R_4 is $-O-R_{11}$, NH_2 , $NHOH$, or is of the formula



R_{11} is hydrogen or C_{1-6} alkyl; and

X is S or O.

40. A compound of the formula:



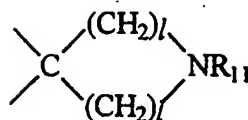
wherein

R_1 may occur one to four times and each occurrence is independently hydrogen or C_{1-6} alkyl;

5 R_2 is hydrogen, pyridyl, C_{1-6} alkyl, $(C_{1-6}$ alkyl)- CO_2-R_{11} , or $-CO_2-R_{11}$;

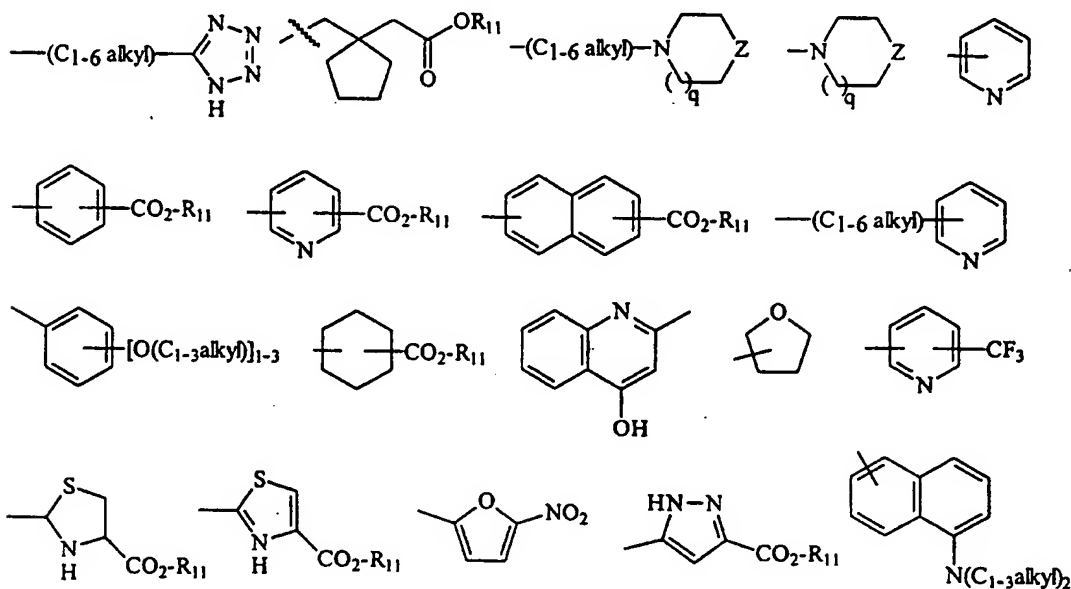
in addition, R_1 and R_2 may be attached to the same carbon atom and form a carbocyclic ring of 5-8 atoms together with the carbon atom to which they are attached, or they may be attached to the same carbon atom and form a ring of 5-8 atoms of the formula:

-300-



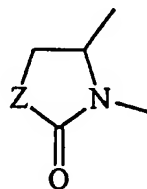
together with the carbon atom to which they are attached;

- 10 R_3 is hydrogen, phenyl, C_{1-6} alkyl, C_{3-6} alkenyl, C_{7-17} arylalkyl, $(C_{1-6}$ alkyl)- CO_2 - R_{11} , $(C_{2-6}$ alkenyl)- CO_2 - R_{11} , $(C_{1-6}$ alkyl)- CO - C_{1-6} alkyl, $(C_{1-6}$ alkyl)- O - C_{1-6} alkyl, $(C_{1-6}$ alkyl)- OH , $(C_{1-6}$ alkyl)- CN , adamantyl or one of the following:

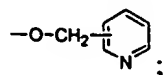


in addition, R_2 and R_3 -(Y)₆ may combine with each other at the terminal thereof to form a ring of the following formula together with the carbon atom and the nitrogen atom

- 15 to which they are attached:

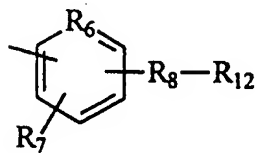


R_4 is $-O-R_{11}$, NH_2 , $NHOH$, $-O-(C_{7-10}$ arylalkyl), or is of the formula



-301-

R_3 is a formula of the following:



;

R_6 is N or CH;

20 R_7 is hydrogen or halogen;

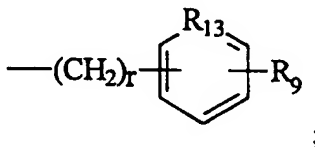
R_8 is $-OCH_2-$;

R_9 may occur one to three times and is a halogen, C_{1-6} alkoxy, C_{1-6} alkyl or trifluoromethyl;

R_{10} is C_{1-6} alkyl, or $(C_{1-6}$ alkyl)-OH, or hydrogen;

25 R_{11} is hydrogen or C_{1-6} alkyl;

R_{12} is C_{1-6} alkyl or the following formula:



;

R_{13} is N or CH;

W is $(C_{1-6}$ alkyl);

30 X is S, O, or CH_2 ;

Y is $-CO-$, $-C(=O)O-$, or $-C(=O)N(R_{10})-$;

Z is O, CH_2 , or $N-R_{11}$;

l is 1, 2, or 3;

m is 1 or 2;

35 n is 0 or 1;

o is 0 or 1;

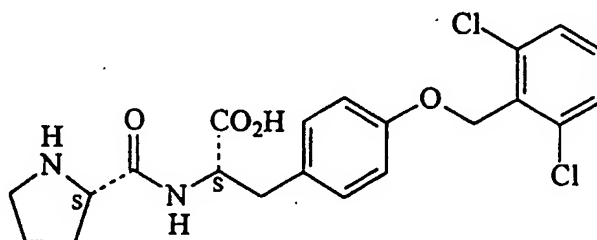
p is 0 or 1;

q is 0 or 1; and

r is 0, 1, 2 or 3;

40 with the provisos that

- (1) when Y is $-C(=O)O-$, R_3 cannot be hydrogen;
- (2) when R_4 is equal to $O-(C_4 \text{ alkyl})$, $C_4 \text{ alkyl}$ is not equal to *tert*-butyl;
- (3) in those pyrrolidine structures (l is 1; m is 2; n is 0; o is 0; p is 1; X is CH_2), W is equal to CH_2 ;
- 45 (4) the compound has an IC_{50} value of less than $5\mu M$ in a Jurkat CS-1 assay and/or an IC_{50} value of less than $50\mu M$ in a Jurkat EC assay;
- (5) when R_3 is phenyl, C_{1-6} alkyl, C_{7-17} arylalkyl, $(C_{1-6} \text{ alkyl})-CO_2-R_{11}$, $(C_{1-6} \text{ alkyl})-O-C_{1-6} \text{ alkyl}$, or $(C_{1-6} \text{ alkyl})OH$, o is 0; and
- (6) the compound is not



INTERNATIONAL SEARCH REPORT

Int. 'onal Application No
PCT/US 99/14233

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07D277/06 A61K31/33 C07D417/12 C07D277/60 C07D281/06
C07D207/16 C07D279/12 C07D417/06 C07D207/26 C07D207/28
C07D211/60 C07D211/62 C07D223/06 C07D513/10 C07D205/04

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC 6 C07D A61K C07K

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	WO 96 22966 A (BIOGEN, INC.) 1 August 1996 (1996-08-01) page 1 -page 2; claim 1 ----	1, 13
A	US 5 688 913 A (THOMAS S. ARRHENIUS ET AL.) 18 November 1997 (1997-11-18) column 1 -column 6 -----	1, 13

☐ Further documents are listed in the continuation of box C.

☒ Patent family members are listed in annex.

* Special categories of cited documents :

- "A" document defining the general state of the art which is not considered to be of particular relevance
- "E" earlier document but published on or after the international filing date
- "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)
- "O" document referring to an oral disclosure, use, exhibition or other means
- "P" document published prior to the international filing date but later than the priority date claimed

- "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
- "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
- "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.
- "Z" document member of the same patent family

Date of the actual completion of the international search

29 September 1999

Date of mailing of the international search report

12/10/1999

Name and mailing address of the ISA

European Patent Office, P.B. 5818 Patentlaan 2
NL - 2280 HV Rijswijk
Tel. (+31-70) 340-2040, Tx. 31 651 epo nl,
Fax: (+31-70) 340-3016

Authorized officer

Van Bijlen, H

INTERNATIONAL SEARCH REPORT

International Application No

PCT/US 99/14233

A. CLASSIFICATION OF SUBJECT MATTER

IPC 6 C07D263/06 C07D401/06 C07D405/06 C07D513/04 C07D417/04
C07D405/04 C07D413/06 C07K5/078 //(C07D513/10,277:00,
221:00),(C07D513/04,277:00,235:00),(C07D513/04,277:00,263:00)

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practical, search terms used)

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.



Further documents are listed in the continuation of box C.



Patent family members are listed in annex.

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"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the International filing date

"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the International filing date but later than the priority date claimed

"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.

"&" document member of the same patent family

Date of the actual completion of the international search

29 September 1999

Date of mailing of the international search report

Name and mailing address of the ISA

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Authorized officer

Van Bijlen, H

INTERNATIONAL SEARCH REPORT

I national application No.

PCT/US 99/ 14233

Box I Observations where certain claims were found unsearchable (Continuation of Item 1 of first sheet)

This International Search Report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☒ Claims Nos.: 14-16
because they relate to subject matter not required to be searched by this Authority, namely:
Remark: Although claims 14-16
are directed to a method of treatment of the human/animal
body, the search has been carried out and based on the alleged
effects of the compound/composition.
2. ☐ Claims Nos.:
because they relate to parts of the International Application that do not comply with the prescribed requirements to such
an extent that no meaningful International Search can be carried out, specifically:
3. ☐ Claims Nos.:
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this International application, as follows:

1. ☐ As all required additional search fees were timely paid by the applicant, this International Search Report covers all
searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment
of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this International Search Report
covers only those claims for which fees were paid, specifically claims Nos.:
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this International Search Report is
restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remark on Protest

☐ The additional search fees were accompanied by the applicant's protest.

☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

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